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(54) Title: NEUROPROTECTIVE PEPTIDES

(57) Abstract: A method of treating a neurodegenerative disorder is disclosed. The method comprises administering to the subject a therapeutically effective amount of an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence no more than 20 amino acids of said CD44V10 amino acid sequence and comprising a neuroprotective activity.

1
NEUROPROTECTIVE PEPTIDES

FIELD AND BACKGROUND OF THE INVENTION

5 The present invention, in some embodiments thereof, relates to neuroprotective peptide agents and uses of same.

10 Neurodegenerative disorders such as Alzheimer's disease (AD), Parkinson's Diseases (PD), Amyotrophic Lateral Sclerosis (ALS) and Huntington's disease (HD), are adult onset, chronic, progressive and irreversible severely disabling diseases in which progressive loss of structure and function of neurons, including death of neurons 15 are present.

15 Alzheimer's disease (AD) is the most prevalent neurodegenerative disorder characterized by progressive loss of cognitive function. AD histopathology is defined by protein abnormalities namely plaques and neurofibrillary tangles which result from deposition of amyloid- β (A β) and hyperphosphorylated tau, respectively. These pathologies are accompanied by loss of neurons and white matter, congophilic angiopathy, inflammation and oxidative damage [1]. The role of inflammation in AD is evidenced by changes in microglia morphology and astrogliosis surrounding the senile plaque [2]. A β peptides are produced from the β -amyloid precursor protein (APP) 20 through an initial β -secretase cleavage followed by the intramembranous digestion by γ -secretase, a protein complex with presenilin1 at its catalytic core. The resulting peptide is secreted and deposited in the AD-defining amyloid plaques [1].

25 Parkinson's disease (PD) is a chronic and progressive neurodegenerative disease caused by a selective degeneration of dopaminergic neurons in the substantia nigra pars compacta of the brain. Symptoms include motor-related, including tremor, rigidity, slowness of movement, and postural instability. Among non-motor symptoms are autonomic dysfunction and sensory and sleep difficulties. Cognitive and neurobehavioral problems, including dementia, are common in the advanced stages of the disease. PD usually appears around the age of 60, although there 30 are young-onset cases. The main pathological characteristic of PD is cell death in the substantia nigra and more specifically the ventral part of the pars compacta, affecting up to 70% of the cells by the time the patient dies [3]

CD44 codes for a family of class I transmembrane proteins which result from extensive alternative splicing and post translation modification. The variations are located in the extracellular membrane-proximal portion of the protein and are encoded by variants exons V2 (V1 in mice) to V10 [4] CD44 is the major cell surface receptor 5 for hyaluronic acid (HA) but it has also been shown to bind proteins such as collagens, fibronectin, fibrinogen, laminin and osteopontin [5]. CD44 is essential for recruitment of circulating lymphocytes to the site of inflammation [6, 7]. CD44S, which doesn't contain any variant exon, is the most ubiquitous form and is expressed by most cell types [8]. CD44 variant proteins, in which one or more of the 10 variant exons are 10 included, are mostly reported in association with cancer [9] and autoimmune diseases such as rheumatoid arthritis [10] and multiple sclerosis [11]. One of the unique functions suggested for CD44 splice variants is participation in signal transduction. As an example it was shown that CD44V6 is essential for signaling through tyrosine kinases such as c-Met [12] and VEGFR-2 [13].

15 In the brain CD44 is found predominantly in astrocytes of the white matter [14-18]. In contrast CD44 variants containing exons V4, V5 and V10 were localized to neurons [17]. CD44 expression was also found in activated microglial cells in the hippocampus following transient forebrain ischemia [19]. CD44 was first mentioned in association with AD when Akiyama et al reported a specific subset of CD44 positive 20 astrocytes which number is increased dramatically in AD brains [14]. CD44 potential role in CNS regeneration was reported as it was found to be essential for axon growth of retinal ganglion cells [20]. CD44 was shown to play a role in ischemic brain injury as CD44-deficient mice had reduced infarct size compared with that of wild-type mice following middle cerebral artery occlusion [21]. Lammich et al [22] reported that CD44 25 goes through dual intramembranous cleavage by a presenilin-dependent secretase [22] that liberates the extracellular domain as well as CD44 intracellular domain for putative nuclear signaling.

WO2009007934 teaches that the expression of splice variants CD44V3, CD44V6 and CD44V10 are significantly increased in the hippocampi of AD patients 30 compared to non-AD individuals

SUMMARY OF THE INVENTION

According to an aspect of some embodiments of the present invention there is provided a method of treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 100 amino acids of the CD44V10 amino acid sequence and comprising a neuroprotective activity, thereby treating the neurodegenerative disorder.

According to an aspect of some embodiments of the present invention there is provided a method of treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid sequence and no more than 100 amino acids of the CD44V6 amino acid sequence and comprising a neuroprotective activity, thereby treating the neurodegenerative disorder.

According to an aspect of some embodiments of the present invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 20 amino acids of the CD44V10 amino acid sequence, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NOS: 49 or 50, the peptide comprising a neuroprotective activity.

According to an aspect of some embodiments of the present invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid sequence and no more than 20 amino acids of the CD44V6 amino acid sequence, the peptide comprising a neuroprotective activity, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NO: 1, 51 or 52.

According to an aspect of some embodiments of the present invention there is provided a pharmaceutical composition comprising as an active agent an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 100 amino acids of a CD44V10 amino acid sequence and comprising a neuroprotective activity and a pharmaceutically effective carrier.

According to an aspect of some embodiments of the present invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 100 amino acids of the CD44V10 amino acid sequence,

and comprising a neuroprotective activity, for use in treating a neurodegenerative disorder.

According to some embodiments of the invention, the peptide comprises an amino acid sequence of formula 1:

5 X₁-G-Y-T-S,

wherein X₁ is any of a glutamic acid or glutamine.

According to some embodiments of the invention, the amino acid sequence comprises peptidomimetics.

According to some embodiments of the invention, the peptidomimetics 10 comprises a retro-inverso mimetic.

According to some embodiments of the invention, the peptide is as set forth in SEQ ID NO: 26, 45 or 46.

According to some embodiments of the invention, the peptide consists of a CD44V10 amino acid sequence.

15 According to some embodiments of the invention, the CD44V10 amino acid sequence is a human CD44V10 amino acid sequence.

According to some embodiments of the invention, the CD44V6 amino acid sequence is a human CD44V6 amino acid sequence.

According to some embodiments of the invention, the peptide comprises a core 20 sequence X₁-X₂-S-H, wherein X₁ and X₂ are acidic amino acids.

According to some embodiments of the invention, X₁ comprises glutamic acid.

According to some embodiments of the invention, X₂ comprises aspartic acid.

According to some embodiments of the invention, the peptide consists of a CD44V6 amino acid sequence.

25 According to some embodiments of the invention, the peptide comprises an amino acid sequence as set forth in SEQ ID NOs: 8-15, 18-45 or 46.

According to some embodiments of the invention, the peptide comprises an amino acid sequence as set forth in SEQ ID NOs: 49 or 50.

According to some embodiments of the invention, the peptide comprises an 30 amino acid sequence as set forth in SEQ ID NOs: 2-7, 16 or 17.

According to an aspect of some embodiments of the present invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid

sequence and no more than 100 amino acids of the CD44V6 amino acid sequence, and comprising a neuroprotective activity, for use in treating a neurodegenerative disorder.

According to some embodiments of the invention, the neurodegenerative disorder is selected from the group consisting of Parkinson's disease, Multiple Sclerosis,

5 ALS, multi-system atrophy, Alzheimer's disease, stroke, traumatic brain injury, progressive supranuclear palsy, fronto-temporal dementia with parkinsonism linked to chromosome 17 and Pick's disease.

According to some embodiments of the invention, the neurodegenerative disease is Parkinson's disease.

10 According to some embodiments of the invention, the neurodegenerative disease is Alzheimer's disease.

According to some embodiments of the invention, the peptide comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 12, 15, 17, 19, 24, 26, 31, 32, 34, 36-38, 43-46.

15 According to some embodiments of the invention, the peptide comprises an amino acid sequence as set forth in SEQ ID NOs: 26 or 45.

According to an aspect of some embodiments of the present invention there is provided a pharmaceutical composition comprising as an active agent an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid sequence and no more than 20 100 amino acids of a CD44V6 amino acid sequence and comprising a neuroprotective activity and a pharmaceutically effective carrier.

According to some embodiments of the invention, the peptide is attached to a cell penetrating agent.

25 According to some embodiments of the invention, at least one of the amino acids is a naturally occurring amino acid.

According to some embodiments of the invention, at least one of the amino acids is a synthetic amino acid.

According to some embodiments of the invention, the synthetic amino acid comprises a D isomer.

30 According to some embodiments of the invention, the isolated peptide is covalently attached to the cell penetrating agent.

According to some embodiments of the invention, the cell penetrating agent is a peptide agent.

According to some embodiments of the invention, the peptide is no longer than 20 amino acids.

5 According to some embodiments of the invention, the peptide is 5-10 amino acids in length.

According to an aspect of some embodiments of the present invention there is provided a method of selecting an agent useful for treating a neurodegenerative disease, the method comprising:

10 (a) contacting a CD44v10/6 peptide with neuronal cells in the presence of a neurotoxic agent; and

(b) monitoring cell death of the neuronal cells, wherein a decrease in an amount or time of cell death of the neuronal cells in the presence of the CD44v10/6 peptide compared to an amount or time of cell death of the neuronal cells in the absence of the
15 CD44v10/6 peptide is indicative of an agent useful for treating a neurodegenerative disease.

According to some embodiments of the invention, the neurotoxic agent is selected from the group consisting of an amyloid peptide, a glutamate, 6-OHDA, MPTP AND MPP+.

20 According to some embodiments of the invention, the administering comprises subcutaneous administering.

According to some embodiments of the invention, the administering comprises intranasal administering.

Unless otherwise defined, all technical and/or scientific terms used herein have
25 the same meaning as commonly understood by one of ordinary skill in the art to which the invention pertains. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of embodiments of the invention, exemplary methods and/or materials are described below. In case of conflict, the patent specification, including definitions, will control. In addition, the materials, methods, and
30 examples are illustrative only and are not intended to be necessarily limiting.

Definitions of the specific embodiments of the invention as claimed herein follow.

According to a first embodiment of the invention, there is provided an isolated peptide comprising:

- (i) at least 5 amino acids of a CD44V10 amino acid sequence and no more than 15 amino acids of said CD44V10 amino acid sequence, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NOs: 49 or 50, the peptide comprising a neuroprotective activity; or
- (ii) at least 3 amino acids of a CD44V6 amino acid sequence and no more than 20 amino acids of said CD44V6 amino acid sequence, the peptide comprising a neuroprotective activity, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NO: 1, 51 or 52,

and wherein said peptide of (i) comprises an amino acid sequence of formula I:

$X_1\text{-G\text{-}Y\text{-}T\text{-}S}$, wherein X_1 is any of a glutamic acid or glutamine;

and wherein said peptide of (ii) comprises a core sequence $X_1\text{-}X_2\text{-S\text{-}H}$, wherein X_1 and X_2 are acidic amino acids.

According to a second embodiment of the invention, there is provided a pharmaceutical composition comprising as an active agent the isolated peptide of the first embodiment and a pharmaceutically effective carrier.

According to a third embodiment of the invention, there is provided a method of treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of the isolated peptide of the first embodiment or the pharmaceutical composition of the second embodiment.

[Text continues on page 7.]

BRIEF DESCRIPTION OF THE DRAWINGS

Some embodiments of the invention are herein described, by way of example only, with reference to the accompanying images. With specific reference now to the drawings in detail, it is stressed that the particulars shown are by way of example and for purposes of illustrative discussion of embodiments of the invention. In this regard, the description taken with the drawings makes apparent to those skilled in the art how embodiments of the invention may be practiced.

In the drawings:

FIGs. 1A-B are multiple protein sequence alignments of CD44V6 (FIG. 1A) and 10 CD44V10 (FIG. 1B) featuring the protein sequences of various mammalian organisms. The alignment was done using protein Blast algorithm online (NCBI). Conserved residues in which there is up to one non-conserved replacement in any of the species are marked by boxes. Non-conserved replacement in conserved residues are marked by red letter.

15 FIG. 2 shows a list of V6 (mouse) and V10 (human) peptides that their synthesis was guided according to the conserved regions denoted in FIG. 1.

FIG. 3: is a bar graph showing the effect of V6 and V10 peptides at 3 concentrations on the viability of SK-N-SH human neuroblastoma cells, following treatment with 80 μ M A β (1-42) for 48 hrs. Cellular viability was determined using the 20 XTT colorimetric assay.

FIGs. 4A-B are bar graphs showing the effect of V6 and V10 peptides at 1 nM on the viability of N2A mouse neuroblastoma cells, following treatment with 25 μ M A β (25-35) for 48 hrs. (FIG. 4A) Relative cellular viability as measured by XTT and (FIG. 4B) relative caspase 3 activity level are shown.

25 FIGs. 5A-B are bar graphs showing the effect of V6 and V10 peptides at 1 μ M on the viability of N2A mouse neuroblastoma cells, following treatment with 200 μ M MPTP for 48 hrs. Relative cellular viability (FIG. 5A) and caspase 3 activity level (FIG. 5B) are shown.

FIGs. 6A-C are graphs showing the effect of human V6 and V10 peptides that 30 are listed in Table 1 and FIG. 2 at 1 pM on the viability of N2A mouse neuroblastoma cells, following treatment with 30 μ M for 48 hrs. Relative viability (FIG. 6A) and caspase 3 activity (FIG. 6B) are shown. (FIG. 6C) The protective effect of N-acetylated

and C-amidates V10A1_N+4 peptide against 30 μ M 6-OHDA in N2A cells was tested at various concentration. Shown is the relative viability as measured by alamar blue fluorescence.

FIGs. 7A-B are bar graphs showing the effect of human V6 and V10 peptides that are listed in Table 1 and FIG. 2 at 1 pM on the viability of SK-N-SH human neuroblastoma cells, following treatment with 25 μ M A β (25-35) for 48 hrs. Relative viability as measured by alamar blue fluorescence (FIG. 7A) and caspase 3 activity (FIG. 7B) are shown.

FIGs. 8A-B is a bar graph showing the effect of human P26-derived peptides that are listed in Table 3 at 10 fM and 1 pM on the viability of N2A mouse neuroblastoma cells, following treatment with 30 μ M 6-OHDA (FIG. 8A) or 40 μ M (FIG. 8B) for 22 hrs. In FIG. 8B the cells were pre-incubated with the peptides for 2.3 hrs prior to the addition of 6-OHDA while in FIG. 8A, 6-OHDA was added together with the peptides. Relative viability as measured by alamar blue fluorescence is shown.

FIG. 9 is a bar graph showing the effects of repeated IH/ICV administration of P26 (SEQ ID NO: 26, 1, 10 and 100 ng/rat) or P34 (SEQ ID NO: 34, 10 and 100 ng/rat) on the discrimination index on the retention test phase in the novel object recognition (NOR) task. ** $P<0.01$ A β ₍₁₋₄₂₎ with vehicle versus the control group (no A β ₍₁₋₄₂₎) and $^{##}P<0.01$ versus the vehicle control.

FIGs. 10A-B are graphs showing the effect of subcutaneous (SC) administered peptides on Morris water maze (MWM) spatial memory assay following A β microinjection. A., The mean latency across 4 training sessions in MWM. Among the peptides-treated groups, the P26 group showed lower levels of the mean latency compared with the vehicle group ($^{##}p<0.001$ in days 2,3 and 4). B. shows the time spent in the target quadrant on test session. The P26 group showed higher levels of the time spent in the target quadrant compared with the vehicle group ($^{##}p<0.05$). UT - untreated group

FIG. 11 is a bar graph showing the effect of SC administered peptides on NOR assay following A β microinjection. The discrimination index on the retention test phase is shown. * $P<0.05$ versus the control group and $^{##}p<0.05$ versus the A β only group

FIG. 12 is a graph showing the pharmacokinetics of P26 (SEQ ID NO: 26) and P26-RI (SEQ ID NO: 45) that was evaluated following subcutaneous administration of

peptide solution in male Sprague Dawley rats at a dose of 1 mg/kg. LC-MS/MS method was used for the quantification of both peptides in plasma samples. The lower limit of quantification (LLOQ) was 22.34 ng/mL.

FIG. 13 is a schematic illustration of the genomic structure of CD44.

5

DESCRIPTION OF SPECIFIC EMBODIMENTS OF THE INVENTION

The present invention, in some embodiments thereof, relates to neuroprotective peptide agents and uses of same.

Before explaining at least one embodiment of the invention in detail, it is to be understood that the invention is not necessarily limited in its application to the details set forth in the following description or exemplified by the Examples. The invention is capable of other embodiments or of being practiced or carried out in various ways.

It was previously found that the expression of splice variants CD44V3, CD44V6 and CD44V10 are significantly increased in the hippocampi of AD patients compared to non-AD individuals. The expression of the CD44 variants was further characterized and found to be mainly neuronal [23].

The present inventors characterized the function of multiple peptides derived from CD44 V6 and V10 exons sequences and found these peptides confer resistance to neuronal cells from neurotoxins such as beta amyloid (A β), MPTP and 6-OHDA, suggesting that these peptides or derivatives may serve as drugs for the treatment of neurodegenerative disorders.

The present inventors have performed structural-functional analyses to uncover minimal active domains that confer neuroprotection. The results were further substantiated in animal models for Parkinson's disease and Alzheimer's disease. These findings place the peptides of the instant invention as lead compounds for drug development.

Thus, according to an aspect of the invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 20 amino acids of said CD44V10 amino acid sequence wherein the peptide comprises a neuroprotective activity.

According to an additional or alternative aspect of the invention there is provided an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid sequence

10

and no more than 20 amino acids of said CD44V6 amino acid sequence, the peptide comprising a neuroprotective activity.

According to one embodiment the CD44V10 amino acid sequence does not consist of the sequence:

5 DSTDRIPATIRNDVTGGRR (SEQ ID NO: 49); or
NSNVNRSLSGDQDTFHPSG (SEQ ID NO: 50).

According to another embodiment the CD44 V6 amino acid sequence does not consist of the sequence:

10 DSTDRIPATIQATPSSTTE (SEQ ID NO: 51); or
DSHSTTGTAGDQDTFHPSG (SEQ ID NO: 52).

Peptides comprising the amino acid sequence set forth in SEQ ID NOS: 49-52 are contemplated for use in the treatment of neurodegenerative diseases, as further elaborated hereinbelow.

15 As used herein "CD44" refers to the cell surface protein that is expressed in a large number of mammalian cell types as set forth in RefSeq Accession No: NM_000610.3. According to a specific embodiment the CD44 is the human CD44 gene. The standard isoform, designated CD44, comprising exons 1-5 and 16-20 is expressed in most cell types. The gene structure is provided in Figure 13 including that of the splice variants CD44V6 and CD44V10.

20 As used herein, the term "CD44V10" corresponds to amino acid coordinates 537-604 of SEQ ID NO: 53, RefSeq Accession No: NP_000601.3 (human CD44 antigen isoform 1 precursor, NCBI Reference Sequence) and is exemplified by SEQ ID NO: 2.

25 As used herein, the term "CD44V6" corresponds to amino acid coordinates 386-427 of SEQ ID NO: 53, RefSeq Accession No: NP_000601.3 (human CD44 antigen isoform 1 precursor, NCBI Reference Sequence) and is exemplified by SEQ ID NO: 8.

30 As used herein, the phrase "neuroprotective activity" refers to prevention of neural cell death. The effect may take the form of protection of neuronal cells i.e., neurons, from apoptosis or degeneration. Assays for qualifying a neuroprotective activity include cell viability assays (e.g., XTT, MTT), morphological assays (e.g., cell staining) or apoptosis biochemical assays (e.g., caspase 3 activity and the like).

According to a specific embodiment, the CD44V6 amino acid sequence is a human CD44V6 amino acid sequence.

According to a specific embodiment, the CD44V10 amino acid sequence is a human CD44V10 amino acid sequence.

5 According to a further specific embodiment, the peptide consists of a CD44V6 amino acid sequence (SEQ ID NO: 2).

10 While further reducing the present invention to practice, the present inventors have uncovered that the peptidic portion (amino acid sequence) which imparts neuroprotection comprises a core sequence X_1-X_2-S-H , wherein X_1 and X_2 are acidic amino acids.

As used herein, the phrase "acidic amino acid" refers to naturally occurring or synthetic amino acids which are polar and negatively charged at physiological pH.

According to a specific embodiment, the X_1 comprises glutamic acid.

According to a specific embodiment, the X_2 comprises aspartic acid.

15 According to a specific embodiment, the peptide comprises the amino acid sequence of SEQ ID NO: 6 or 7.

According to a further embodiment, the CD44V6 consists of a CD44V6 amino acid sequence (SEQ ID NO: 2).

20 According to a further embodiment, the peptide comprises an amino acid sequence as set forth in SEQ ID NOs: 2-7, 16 or 17.

As mentioned, peptides of CD44V10 are also contemplated herein. Thus, according to an exemplary embodiment the peptide comprises an amino acid sequence as set forth in SEQ ID NOs: 8-15, 18-46, or specifically, SEQ ID NO: 8-15, 18-38, 39-42 or 43-46.

25 While further reducing the present invention to practice, the present inventors were able to identify a minimal portion of CD44V10 which is active in conferring neuroprotection.

Thus, according to an exemplary embodiment, the CD44V10 peptide comprises an amino acid sequence of formula 1:

30 $X_1-G-Y-T-S$,

wherein X_1 is any of a glutamic acid or glutamine.

As will be further described in details hereinbelow, the amino acid sequence of the peptide comprises peptidomimetics, such as a retro-inverso mimetic (e.g., SEQ ID NO: 45 or 46).

According to an exemplary embodiment, the peptide is as set forth in SEQ ID NO: 26.

According to an exemplary embodiment, the peptide is as set forth in SEQ ID NO: 26, 45 or 46.

According to a further specific embodiment, the peptide consists of a CD44V10 amino acid sequence (SEQ ID NO: 8).

The term "peptide" as used herein refers to a polymer of natural or synthetic amino acids, encompassing native peptides (either degradation products, synthetically synthesized polypeptides or recombinant polypeptides) and peptidomimetics (e.g., inverso, retro or retro-inverso, typically, synthetically synthesized peptides), as well as peptoids and semipeptoids which are polypeptide analogs, which may have, for example, modifications rendering the peptides even more stable while in a body or more capable of penetrating into cells.

Such modifications include, but are not limited to N terminus modification, C terminus modification, polypeptide bond modification, including, but not limited to, CH₂-NH, CH₂-S, CH₂-S=O, O=C-NH, CH₂-O, CH₂-CH₂, S=C-NH, CH=CH or CF=CH, backbone modifications, and residue modification. Methods for preparing peptidomimetic compounds are well known in the art and are specified, for example, in Quantitative Drug Design, C.A. Ramsden Gd., Chapter 17.2, F. Choplin Pergamon Press (1992), which is incorporated by reference as if fully set forth herein. Further details in this respect are provided hereinunder.

Polypeptide bonds (-CO-NH-) within the polypeptide may be substituted, for example, by N-methylated bonds (-N(CH₃)-CO-), ester bonds (-C(R)H-C-O-O-C(R)-N-), ketomethylen bonds (-CO-CH₂-), α -aza bonds (-NH-N(R)-CO-), wherein R is any alkyl, e.g., methyl, carba bonds (-CH₂-NH-), hydroxyethylene bonds (-CH(OH)-CH₂-), thioamide bonds (-CS-NH-), olefinic double bonds (-CH=CH-), retro amide bonds (-NH-CO-), polypeptide derivatives (-N(R)-CH₂-CO-), wherein R is the "normal" side chain, naturally presented on the carbon atom.

These modifications can occur at any of the bonds along the polypeptide chain and even at several (2-3) at the same time.

Natural aromatic amino acids, Trp, Tyr and Phe, may be substituted for synthetic non-natural acid such as Phenylglycine, naphthylealanine (Nol), ring-methylated derivatives of Phe, halogenated derivatives of Phe or o-methyl-Tyr.

In addition to the above, the polypeptides of the present invention may also include one or more modified amino acids or one or more non-amino acid monomers (e.g. fatty acids, complex carbohydrates etc).

As used herein in the specification and in the claims section below the term "amino acid" or "amino acids" is understood to include the 20 naturally occurring amino acids; those amino acids often modified post-translationally *in vivo*, including, for example, hydroxyproline, phosphoserine and phosphothreonine; and other unusual amino acids including, but not limited to, 2-amino adipic acid, hydroxylysine, isodesmosine, nor-valine, nor-leucine and ornithine. Furthermore, the term "amino acid" includes both D- and L-amino acids (stereoisomers).

Tables A and B below list naturally occurring amino acids (Table A) and non-conventional or modified amino acids (Table B) which can be used with the present invention.

Table A

Amino Acid	Three-Letter Abbreviation	One-letter Symbol
alanine	Ala	A
Arginine	Arg	R
Asparagine	Asn	N
Aspartic acid	Asp	D
Cysteine	Cys	C
Glutamine	Gln	Q
Glutamic Acid	Glu	E
glycine	Gly	G
Histidine	His	H
isoleucine	Ile	I
leucine	Leu	L
Lysine	Lys	K
Methionine	Met	M
phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S

Amino Acid	Three-Letter Abbreviation	One-letter Symbol
Threonine	Thr	T
tryptophan	Trp	W
tyrosine	Tyr	Y
Valine	Val	V
Any amino acid as above	Xaa	X

Table B

Non-conventional amino acid	Code	Non-conventional amino acid	Code
α -aminobutyric acid	Abu	L-N-methylalanine	Nmala
α -amino- α - methylbutyrate	Mgabu	L-N-methylarginine	Nmarg
aminocyclopropane-	Cpro	L-N-methylasparagine	Nmasn
carboxylate		L-N-methylaspartic acid	Nmasp
aminoisobutyric acid	Aib	L-N-methylcysteine	Nmcys
aminonorbornyl-	Norb	L-N-methylglutamine	Nmgin
carboxylate		L-N-methylglutamic acid	Nmglu
cyclohexylalanine	Chexa	L-N-methylhistidine	Nmhis
cyclopentylalanine	Cpen	L-N-methylisoleucine	Nmile
D-alanine	Dal	L-N-methylleucine	Nmleu
D-arginine	Darg	L-N-methyllysine	Nmlys
D-aspartic acid	Dasp	L-N-methylmethionine	Nmmet
D-cysteine	Dcys	L-N-methylnorleucine	Nmnle
D-glutamine	Dgln	L-N-methylnorvaline	Nmnva
D-glutamic acid	Dglu	L-N-methylornithine	Nmorn
D-histidine	Dhis	L-N-methylphenylalanine	Nmphe
D-isoleucine	Dile	L-N-methylproline	Nmpro
D-leucine	Dleu	L-N-methylserine	Nmser
D-lysine	Dlys	L-N-methylthreonine	Nmthr
D-methionine	Dmet	L-N-methyltryptophan	Nmtrp
D-ornithine	Dorn	L-N-methyltyrosine	Nmtyr
D-phenylalanine	Dphe	L-N-methylvaline	Nmval
D-proline	Dpro	L-N-methylethylglycine	Nmetg

D-serine	Dser	L-N-methyl-t-butylglycine	Nmtbug
D-threonine	Dthr	L-norleucine	Nle
D-tryptophan	Dtrp	L-norvaline	Nva
D-tyrosine	Dtyr	α -methyl-aminoisobutyrate	Maib
D-valine	Dval	α -methyl- γ -aminobutyrate	Mgabu
D- α -methylalanine	Dmala	α ethylcyclohexylalanine	Mchexa
D- α -methylarginine	Dmarg	α methylcyclopentylalanine	Mcpen
D- α -methylasparagine	Dmasn	α -methyl- α -naphthylalanine	Manap
D- α -methylaspartate	Dmasp	α methylpenicillamine	Mpen
D- α -methylcysteine	Dmcys	N-(4-aminobutyl)glycine	Nglu
D- α -methylglutamine	Dmgln	N-(2-aminoethyl)glycine	Naeg
D- α -methylhistidine	Dmhis	N-(3-aminopropyl)glycine	Norn
D- α -methylisoleucine	Dmile	N- amino- α -methylbutyrate	Nmaabu
D- α -methylleucine	Dmleu	α -naphthylalanine	Anap
D- α -methyllysine	Dmlys	N-benzylglycine	Nphe
D- α -methylmethionine	Dmmet	N-(2-carbamylethyl)glycine	Ngln
D- α -methylornithine	Dmorn	N- (carbamylmethyl)glycine	Nasn
D- α -methylphenylalanine	Dmphe	N-(2-carboxyethyl)glycine	Nglu
D- α -methylproline	Dmpro	N- (carboxymethyl)glycine	Nasp
D- α -methylserine	Dmser	N-cyclobutylglycine	Ncbut
D- α -methylthreonine	Dmthr	N- cycloheptylglycine	Nchep
D- α -methyltryptophan	Dmtrp	N-cyclohexylglycine	Nchex
D- α -methyltyrosine	Dmty	N-cyclodecylglycine	Ncdec
D- α -methylvaline	Dmval	N- cyclododecylglycine	Ncdod
D- α -methylalanine	Dnmala	N-cyclooctylglycine	Ncoct
D- α -methylarginine	Dnmarg	N- cyclopropylglycine	Ncpro
D- α -methylasparagine	Dnmasn	N- cycloundecylglycine	Ncund

D- α -methylasparatate	Dnmasp	N-(2,2-diphenylethyl)glycine	Nbhm
D- α -methylcysteine	Dnmcys	N-(3,3-diphenylpropyl)glycine	Nbhe
D-N-methylleucine	Dnmleu	N-(3-indolyethyl)glycine	Nhtrp
D-N-methyllysine	Dnmlys	N-methyl- γ -aminobutyrate	Nmgabu
N-methylcyclohexylalanine	Nmchex	D-N-methylmethionine	Dnmme
D-N-methylornithine	Dnmorn	N-methylcyclopentylalanine	Nmcpen
N-methylglycine	Nala	D-N-methylphenylalanine	Dnmphe
N-methylaminoisobutyrate	Nmaib	D-N-methylproline	Dnmpro
N-(1-methylpropyl)glycine	Nile	D-N-methylserine	Dnmser
N-(2-methylpropyl)glycine	Nile	D-N-methylserine	Dnmser
N-(2-methylpropyl)glycine	Nleu	D-N-methylthreonine	Dnmthr
D-N-methyltryptophan	Dnmtrp	N-(1-methylethyl)glycine	Nva
D-N-methyltyrosine	Dnmtyr	N-methyl- α -naphthylalanine	Nmanap
D-N-methylvaline	Dnmval	N-methylpenicillamine	Nmpen
γ -aminobutyric acid	Gaba	N-(<i>p</i> -hydroxyphenyl)glycine	Nhtyr
L- <i>t</i> -butylglycine	Tbug	N-(thiomethyl)glycine	Ncys
L-ethylglycine	Etg	penicillamine	Pen
L-homophenylalanine	Hphe	L- α -methylalanine	Mala
L- α -methylarginine	Marg	L- α -methylasparagine	Masn
L- α -methylaspartate	Masp	L- α -methyl- <i>t</i> -butylglycine	Mtbug
L- α -methylcysteine	Mcys	L-methylethylglycine	Metg
L- α -thylglutamine	Mgln	L- α -methylglutamate	Mglu
L- α -methylhistidine	Mhis	L- α -methylhomo-phenylalanine	Mhphe
L- α -methylisoleucine	Mile	N-(2-methylthioethyl)glycine	Nmet

D-N-methylglutamine	Dnmgln	N-(3-guanidinopropyl)glycine	Narg
D-N-methylglutamate	Dnmglu	N-(1-hydroxyethyl)glycine	Nthr
D-N-methylhistidine	Dnmhis	N-(hydroxyethyl)glycine	Nser
D-N-methylisoleucine	Dnmile	N-(imidazolylethyl)glycine	Nhis
D-N-methylleucine	Dnmleu	N-(3-indolylethyl)glycine	Nhtrp
D-N-methyllysine	Dnmlys	N-methyl- γ -aminobutyrate	Nmgabu
N-methylcyclohexylalanine	Nmchex	D-N-methylmethionine	Dnmme
D-N-methylornithine	Dnmorn	N-methylcyclopentylalanine	Nmcpen
N-methylglycine	Nala	D-N-methylphenylalanine	Dnmphe
N-methylaminoisobutyrate	Nmaib	D-N-methylproline	Dnmpro
N-(1-methylpropyl)glycine	Nile	D-N-methylserine	Dnmser
N-(2-methylpropyl)glycine	Nleu	D-N-methylthreonine	Dnmthr
D-N-methyltryptophan	Dnmtrp	N-(1-methylethyl)glycine	Nval
D-N-methyltyrosine	Dnmtyr	N-methyl- α -naphthylalanine	Nmanap
D-N-methylvaline	Dnmval	N-methylpenicillamine	Nmpen
γ -aminobutyric acid	Gabu	N-(<i>p</i> -hydroxyphenyl)glycine	Nhtyr
L- <i>t</i> -butylglycine	Tbug	N-(thiomethyl)glycine	Ncys
L-ethylglycine	Etg	penicillamine	Pen
L-homophenylalanine	Hphe	L- α -methylalanine	Mala
L- α -methylarginine	Marg	L- α -methylasparagine	Masn
L- α -methylaspartate	Masp	L- α -methyl- <i>t</i> -butylglycine	Mtbug
L- α -methylcysteine	Mcys	L- α -methylethylglycine	Metg
L- α -methylglutamine	Mgln	L- α -methylglutamate	Mglu
L- α -ethylhistidine	Mhis	L- α -methylhomophenylalanine	Mhphe

L- α thylisoleucine	Mile	N-(2-methylthioethyl)glycine	Nmet
L- α -methylleucine	Mleu	L- α -methyllysine	Mlys
L- α - methylmethionine	Mmet	L- α - methylnorleucine	Mnle
L- α - methylnorvaline	Mnva	L- α - methylornithine	Morn
L- α - methylphenylalanine	Mphe	L- α - methylproline	Mpro
L- α - methylserine	mser	L- α - methylthreonine	Mthr
L- α ethylvaline	Mtrp	L- α - methyltyrosine	Mtyr
L- α -methylleucine	Mval Nnbhm	L-N- methylhomophenylalanine	Nmhphe
N-(N-(2,2-diphenylethyl) carbamylmethyl-glycine	Nnbhm	N-(N-(3,3-diphenylpropyl) carbamylmethyl(1)glycine	Nnbhe
1-carboxy-1-(2,2-diphenyl ethylamino)cyclopropane	Nmbc		

Table B Cont.

The amino acids of the peptides of the present invention may be substituted either conservatively or non-conservatively.

The term “conservative substitution” as used herein, refers to the replacement of 5 an amino acid present in the native sequence in the peptide with a naturally or non-naturally occurring amino or a peptidomimetics having similar steric properties. Where the side-chain of the native amino acid to be replaced is either polar or hydrophobic, the conservative substitution should be with a naturally occurring amino acid, a non-naturally occurring amino acid or with a peptidomimetic moiety which is also polar or 10 hydrophobic (in addition to having the same steric properties as the side-chain of the replaced amino acid).

As naturally occurring amino acids are typically grouped according to their properties, conservative substitutions by naturally occurring amino acids can be easily determined bearing in mind the fact that in accordance with the invention replacement 15 of charged amino acids by sterically similar non-charged amino acids are considered as conservative substitutions.

For producing conservative substitutions by non-naturally occurring amino acids it is also possible to use amino acid analogs (synthetic amino acids) well known in the

art. A peptidomimetic of the naturally occurring amino acid is well documented in the literature known to the skilled practitioner.

When affecting conservative substitutions the substituting amino acid should have the same or a similar functional group in the side chain as the original amino acid.

5 The phrase "non-conservative substitutions" as used herein refers to replacement of the amino acid as present in the parent sequence by another naturally or non-naturally occurring amino acid, having different electrochemical and/or steric properties. Thus, the side chain of the substituting amino acid can be significantly larger (or smaller) than the side chain of the native amino acid being substituted and/or can have functional 10 groups with significantly different electronic properties than the amino acid being substituted. Examples of non-conservative substitutions of this type include the substitution of phenylalanine or cyclohexylmethyl glycine for alanine, isoleucine for glycine, or -NH-CH[(-CH₂)₅-COOH]-CO- for aspartic acid. Those non-conservative 15 substitutions which fall under the scope of the present invention are those which still constitute a peptide having neuroprotective properties.

As mentioned, the N and C termini of the peptides of the present invention may be protected by function groups. Suitable functional groups are described in Green and Wuts, "Protecting Groups in Organic Synthesis", John Wiley and Sons, Chapters 5 and 7, 1991, the teachings of which are incorporated herein by reference. Preferred protecting groups 20 are those that facilitate transport of the compound attached thereto into a cell, for example, by reducing the hydrophilicity and increasing the lipophilicity of the compounds.

These moieties can be cleaved *in vivo*, either by hydrolysis or enzymatically, inside the cell. Hydroxyl protecting groups include esters, carbonates and carbamate protecting groups. Amine protecting groups include alkoxy and aryloxy carbonyl groups, as 25 described above for N-terminal protecting groups. Carboxylic acid protecting groups include aliphatic, benzylic and aryl esters, as described above for C-terminal protecting groups. In one embodiment, the carboxylic acid group in the side chain of one or more glutamic acid or aspartic acid residue in a peptide of the present invention is protected, preferably with a methyl, ethyl, benzyl or substituted benzyl ester.

30 Examples of N-terminal protecting groups include acyl groups (-CO-R1) and alkoxy carbonyl or aryloxy carbonyl groups (-CO-O-R1), wherein R1 is an aliphatic, substituted aliphatic, benzyl, substituted benzyl, aromatic or a substituted aromatic group.

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Specific examples of acyl groups include acetyl, (ethyl)-CO-, n-propyl-CO-, iso-propyl-CO-, n-butyl-CO-, sec-butyl-CO-, t-butyl-CO-, hexyl, lauroyl, palmitoyl, myristoyl, stearyl, oleoyl phenyl-CO-, substituted phenyl-CO-, benzyl-CO- and (substituted benzyl)-CO-. Examples of alkoxy carbonyl and aryloxy carbonyl groups 5 include CH₃-O-CO-, (ethyl)-O-CO-, n-propyl-O-CO-, iso-propyl-O-CO-, n-butyl-O-CO-, sec-butyl-O-CO-, t-butyl-O-CO-, phenyl-O-CO-, substituted phenyl-O-CO- and benzyl-O-CO-, (substituted benzyl)-O-CO-. Adamantan, naphtalen, myristoleyl, tuluen, biphenyl, cinnamoyl, nitrobenzoy, toluoyl, furoyl, benzoyl, cyclohexane, norbornane, Z-caproic. In order to facilitate the N-acylation, one to four glycine residues can be present 10 in the N-terminus of the molecule.

The carboxyl group at the C-terminus of the compound can be protected, for example, by an amide (i.e., the hydroxyl group at the C-terminus is replaced with -NH₂, -NHR₂ and -NR₂R₃) or ester (i.e. the hydroxyl group at the C-terminus is replaced with -OR₂). R₂ and R₃ are independently an aliphatic, substituted aliphatic, benzyl, substituted 15 benzyl, aryl or a substituted aryl group. In addition, taken together with the nitrogen atom, R₂ and R₃ can form a C4 to C8 heterocyclic ring with from about 0-2 additional heteroatoms such as nitrogen, oxygen or sulfur. Examples of suitable heterocyclic rings include piperidinyl, pyrrolidinyl, morpholino, thiomorpholino or piperazinyl. Examples of C-terminal protecting groups include -NH₂, -NHCH₃, -N(CH₃)₂, -NH(ethyl), -N(ethyl)₂, 20 -N(methyl) (ethyl), -NH(benzyl), -N(C1-C4 alkyl)(benzyl), -NH(phenyl), -N(C1-C4 alkyl) (phenyl), -OCH₃, -O-(ethyl), -O-(n-propyl), -O-(n-butyl), -O-(iso-propyl), -O-(sec- butyl), -O-(t-butyl), -O-benzyl and -O-phenyl.

Of note, peptides of the invention (derived from either CD44V6, CD44V10, as described above) are referred to in general as CD44 peptides of the invention.

25 The CD44 peptides of the invention (i.e., the neuroprotecting peptide portion) is 3-100, or 3-50, or 3-40, or 3-30 amino acids in length. According to further embodiments, the peptide is 3-20, 5-20, 5-20, 5-18, 5-15, 5-10, 7-10, 8-10 amino acids in length.

The CD44 peptides of the invention may be qualified for their neuroprotective activity as described hereinabove and in the Examples section which follows using both in 30 vitro and in vivo models for neuroprotection and neurodegenerative conditions.

The present teachings may further be employed for the identification of agents useful for treating a neurodegenerative disease.

Thus, there is provided a method comprising:

5 (a) contacting a CD44v10/6 peptide with neuronal cells in the presence of a neurotoxic agent; and

10 (b) monitoring cell death of said neuronal cells, wherein a decrease in an amount or time of cell death of said neuronal cells in the presence of said CD44v10/6 peptide compared to an amount or time of cell death of said neuronal cells in the absence of said CD44v10/6 peptide is indicative of an agent useful for treating a neurodegenerative disease.

Methods of monitoring neural cell death are well known in the art and are further described hereinabove (under neuroprotection) and in the Examples section which follows.

15 A neurotoxic agent as used herein refers to a molecule a condition or state that damages the nervous system and/or brain, usually by killing neurons.

According to a specific embodiment, the neurotoxic agent is selected from the group consisting of an amyloid, a glutamate, 6-OHDA, MPTP AND MPP⁺.

20 In order to improve the bioavailability of the CD44 peptides, a single, a portion or even all the amino acids in the peptide can be D amino acids which are not susceptible to enzymatic proteolytic activity and can improve altogether the use of the peptides of the invention as pharmaceuticals. The peptides of the present invention may be attached (either covalently or non-covalently) to a penetrating agent.

As used herein the phrase "penetrating agent" refers to an agent which enhances translocation of any of the attached peptide across a cell membrane.

25 According to one embodiment, the penetrating agent is a peptide and is attached to the CD44 peptides (either directly or non-directly) via a peptide bond.

30 Typically, peptide penetrating agents have an amino acid composition containing either a high relative abundance of positively charged amino acids such as lysine or arginine, or have sequences that contain an alternating pattern of polar/charged amino acids and non-polar, hydrophobic amino acids.

By way of non-limiting example, cell penetrating peptide (CPP) sequences may be used in order to enhance intracellular penetration. CPPs may include short and long

versions of the protein transduction domain (PTD) of HIV TAT protein [YGRKKRR (SEQ ID NO: 54), YGRKKRRQRR (SEQ ID NO: 55), or RRQRR (SEQ ID NO: 56)]. However, the disclosure is not so limited, and any suitable penetrating agent may be used, as known by those of skill in the art.

5 According to a particular embodiment, the peptide conjugates of the present invention are no longer than 25, 30 or 40 amino acids (this includes the CD44 peptide together with any additional attached sequence, such as a cell penetrating peptide as described above).

10 The peptides of the present invention may also comprise non-amino acid moieties, such as for example, hydrophobic moieties (various linear, branched, cyclic, polycyclic or heterocyclic hydrocarbons and hydrocarbon derivatives) attached to the peptides; non-peptide penetrating agents; various protecting groups, especially where the compound is linear, which are attached to the compound's terminals to decrease degradation. Chemical (non-amino acid) groups present in the compound may be included in order to improve 15 various physiological properties such; decreased degradation or clearance; decreased repulsion by various cellular pumps, improve immunogenic activities, improve various modes of administration (such as attachment of various sequences which allow penetration through various barriers, through the gut, etc.); increased specificity, increased affinity, decreased toxicity and the like.

20 Attaching the amino acid sequence component of the peptides of the invention to other non-amino acid agents may be by covalent linking, by non-covalent complexion, for example, by complexion to a hydrophobic polymer, which can be degraded or cleaved producing a compound capable of sustained release; by entrapping the amino acid part of the peptide in liposomes or micelles to produce the final peptide of the invention. The 25 association may be by the entrapment of the amino acid sequence within the other component (liposome, micelle) or the impregnation of the amino acid sequence within a polymer to produce the final peptide of the invention.

30 The peptides of the invention may be linear or cyclic (cyclization may improve stability). Cyclization may take place by any means known in the art. Where the compound is composed predominantly of amino acids, cyclization may be via N- to C-terminal, N-terminal to side chain and N-terminal to backbone, C-terminal to side chain, C-terminal to backbone, side chain to backbone and side chain to side chain, as well as

backbone to backbone cyclization. Cyclization of the peptide may also take place through non-amino acid organic moieties comprised in the peptide.

The peptides of the present invention can be biochemically synthesized such as by using standard solid phase techniques. These methods include exclusive solid phase synthesis, partial solid phase synthesis methods, fragment condensation, classical solution synthesis. Solid phase polypeptide synthesis procedures are well known in the art and further described by John Morrow Stewart and Janis Dillaha Young, Solid Phase Polypeptide Syntheses (2nd Ed., Pierce Chemical Company, 1984).

Large scale peptide synthesis is described by Andersson Biopolymers 10 2000;55(3):227-50.

Synthetic peptides can be purified by preparative high performance liquid chromatography [Creighton T. (1983) Proteins, structures and molecular principles. WH Freeman and Co. N.Y.] and the composition of which can be confirmed via amino acid sequencing.

15 Recombinant techniques may also be used to generate the peptides of the present invention. To produce a peptide of the present invention using recombinant technology, a polynucleotide encoding the peptide of the present invention is ligated into a nucleic acid expression vector, which comprises the polynucleotide sequence under the transcriptional control of a *cis*-regulatory sequence (e.g., promoter sequence) suitable 20 for directing constitutive, tissue specific or inducible transcription of the polypeptides of the present invention in the host cells.

In addition to being synthesizable in host cells, the peptides of the present invention can also be synthesized using *in vitro* expression systems. These methods are well known in the art and the components of the system are commercially available.

25 As mentioned, by virtue of their neuroprotective function, the peptides of the present invention may be used to treat neurodegenerative disorders.

Thus according to an aspect of the invention, there is provided a method of 30 treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of an isolated peptide comprising at least 3 amino acids of a CD44V10 amino acid sequence and no more than 20 amino acids of said CD44V10 amino acid sequence and comprising a neuroprotective activity, thereby treating the neurodegenerative disorder.

According to another aspect of the invention, there is provided a method of treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of an isolated peptide comprising at least 3 amino acids of a CD44V6 amino acid sequence and no more than 5 20 amino acids of said CD44V6 amino acid sequence and comprising a neuroprotective activity, thereby treating the neurodegenerative disorder.

As used herein, the phrase "a subject in need thereof" or "a subject" refers to mammals, preferably human beings at any age which suffer from a neural damage or is at risk to develop a neural damage.

10 As used herein the phrase "neural damage" refers to any disease, disorder or condition which is characterized by an acute and/or progressive damage and/or loss of neuronal cells and/or glial cells.

According to some embodiments of the invention, the pathology associated with neural damage affects neuronal and/or glial cells in the central nervous system.

15 Non-limiting examples of pathologies caused by an acute or sudden damage to neuronal cells include brain injury, spinal injury, head injury, and stroke [cerebrovascular accident (CVA)].

20 According to some embodiments of the invention, the pathology associated with neural damage is cancer. Non-limiting examples of cancers which affect the neuronal and glial cells include glioblastoma, neuroblastoma, adenocarcinoma of the brain, as well as metastases of a distant cancers such as breast cancer, lung cancer, and the like.

According to some embodiments of the invention, the pathology associated with neural damage is chronic.

25 According to some embodiments of the invention, the pathology associated with neural damage is a neurodegenerative disease.

Exemplary neurodegenerative diseases or conditions include, but are not limited to multi-system atrophy, stroke, progressive supranuclear palsy, fronto-temporal dementia with parkinsonism linked to chromosome 17, traumatic brain injury (TBI), Pick's disease, multiple sclerosis, Lupus eruthromatosis, Alzheimer's disease, 30 Parkinson's Disease, senile dementia, amyotrophic lateral sclerosis, Down's Syndrome, Dutch Type Hereditary Cerebral Hemorrhage Amyloidosis, Reactive Amyloidosis, Familial Mediterranean Fever, Familial Amyloid Nephropathy with Urticaria and

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Deafness, Muckle-Wells Syndrome, Idiopathic Myeloma, Macroglobulinemia-Associated Myeloma, Familial Amyloid Polyneuropathy, Familial Amyloid Cardiomyopathy, Isolated Cardiac Amyloid, Systemic Senile Amyloidosis, Adult Onset Diabetes, Insulinoma, Isolated Atrial Amyloid, Medullary Carcinoma of the Thyroid, 5 Familial Amyloidosis, Hereditary Cerebral Hemorrhage with Amyloidosis, Familial Amyloidotic Polyneuropathy, Scrapie, Creutzfeldt-Jacob Disease, Gerstmann Straussler-Scheinker Syndrome, Bovine Spongiform Encephalitis, a Prion-mediated disease, and Huntington's Disease.

According to a specific embodiment, the neurodegenerative disease is 10 Alzheimer's disease.

According to a specific embodiment, the neurodegenerative disease is Parkinson's disease.

The peptides of the present invention may be provided *per se* or as part of a pharmaceutical composition, where it is mixed with suitable carriers or excipients.

15 As used herein a "pharmaceutical composition" refers to a preparation of one or more of the active ingredients described herein with other chemical components such as physiologically suitable carriers and excipients. The purpose of a pharmaceutical composition is to facilitate administration of a compound to an organism.

Herein the term "active ingredient" refers to the peptides accountable for the 20 biological effect.

Hereinafter, the phrases "physiologically acceptable carrier" and "pharmaceutically acceptable carrier" which may be interchangeably used refer to a carrier or a diluent that does not cause significant irritation to an organism and does not abrogate the biological activity and properties of the administered compound. An 25 adjuvant is included under these phrases.

Herein the term "excipient" refers to an inert substance added to a pharmaceutical composition to further facilitate administration of an active ingredient. Examples, without limitation, of excipients include calcium carbonate, calcium phosphate, various sugars and types of starch, cellulose derivatives, gelatin, vegetable 30 oils and polyethylene glycols.

Techniques for formulation and administration of drugs may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, PA, latest edition, which is incorporated herein by reference.

Suitable routes of administration may, for example, include oral, rectal, transmucosal, especially transnasal, intestinal or parenteral delivery, including intramuscular, subcutaneous and intramedullary injections as well as intrathecal, direct intraventricular, intracardiac, e.g., into the right or left ventricular cavity, into the common coronary artery, intravenous, intraperitoneal, intranasal, or intraocular injections.

As described in length in the Examples section which follows, the peptides of the invention were able to protect against Alzheimer's and Parkinson's disease when administered directly into the brain such as by intrahippocampal (IH) intracerebroventricular injection (ICV), intracranial (IC) or intrathecal administration, essentially providing for a local mode of administration, each of which is contemplated herein.

Conventional approaches for drug delivery to the central nervous system (CNS) include: neurosurgical strategies (e.g., intracerebral injection or intracerebroventricular infusion); molecular manipulation of the agent (e.g., production of a chimeric fusion protein that comprises a transport peptide that has an affinity for an endothelial cell surface molecule in combination with an agent that is itself incapable of crossing the BBB) in an attempt to exploit one of the endogenous transport pathways of the BBB; pharmacological strategies designed to increase the lipid solubility of an agent (e.g., conjugation of water-soluble agents to lipid or cholesterol carriers); and the transitory disruption of the integrity of the BBB by hyperosmotic disruption (resulting from the infusion of a mannitol solution into the carotid artery or the use of a biologically active agent such as an angiotensin peptide). However, each of these strategies has limitations, such as the inherent risks associated with an invasive surgical procedure, a size limitation imposed by a limitation inherent in the endogenous transport systems, potentially undesirable biological side effects associated with the systemic administration of a chimeric molecule comprised of a carrier motif that could be active outside of the CNS, and the possible risk of brain damage within regions of the brain where the BBB is disrupted, which renders it a suboptimal delivery method.

Pharmaceutical compositions of the present invention may be manufactured by processes well known in the art, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes.

5 Pharmaceutical compositions for use in accordance with the present invention thus may be formulated in conventional manner using one or more physiologically acceptable carriers comprising excipients and auxiliaries, which facilitate processing of the active ingredients into preparations which, can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

10 For injection, the active ingredients of the pharmaceutical composition may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hank's solution, Ringer's solution, or physiological salt buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

15 For oral administration, the pharmaceutical composition can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well known in the art. Such carriers enable the pharmaceutical composition to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions, and the like, for oral ingestion by a patient. Pharmacological preparations for oral use
20 can be made using a solid excipient, optionally grinding the resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth,
25 methyl cellulose, hydroxypropylmethyl-cellulose, sodium carbomethylcellulose; and/or physiologically acceptable polymers such as polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

30 Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, titanium dioxide, lacquer solutions and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to

the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical compositions which can be used orally, include push-fit capsules made of gelatin as well as soft, sealed capsules made of gelatin and a plasticizer, such as 5 glycerol or sorbitol. The push-fit capsules may contain the active ingredients in admixture with filler such as lactose, binders such as starches, lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active ingredients may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added. All formulations for 10 oral administration should be in dosages suitable for the chosen route of administration.

For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by nasal inhalation, the active ingredients for use according to the present invention are conveniently delivered in the form of an aerosol spray 15 presentation from a pressurized pack or a nebulizer with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichloro-tetrafluoroethane or carbon dioxide. In the case of a pressurized aerosol, the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., 20 gelatin for use in a dispenser may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

The pharmaceutical composition described herein may be formulated for parenteral administration, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampoules or in multidose containers with optionally, an added preservative. The compositions may be 25 suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

Pharmaceutical compositions for parenteral administration include aqueous solutions of the active preparation in water-soluble form. Additionally, suspensions of the active ingredients may be prepared as appropriate oily or water based injection 30 suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acids esters such as ethyl oleate, triglycerides or liposomes. Aqueous injection suspensions may contain substances, which increase the viscosity of

the suspension, such as sodium carboxymethyl cellulose, sorbitol or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the active ingredients to allow for the preparation of highly concentrated solutions.

5 Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile, pyrogen-free water based solution, before use.

The pharmaceutical composition of the present invention may also be formulated in rectal compositions such as suppositories or retention enemas, using, e.g., conventional suppository bases such as cocoa butter or other glycerides.

10 Pharmaceutical compositions suitable for use in context of the present invention include compositions wherein the active ingredients are contained in an amount effective to achieve the intended purpose. More specifically, a therapeutically effective amount means an amount of active ingredients (CD44 peptides) effective to prevent, alleviate or ameliorate symptoms of a disorder (e.g., Parkinson's Disease, Alzheimer's disease) or prolong the survival of the subject being treated.

15 Determination of a therapeutically effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein.

For any preparation used in the methods of the invention, the therapeutically effective amount or dose can be estimated initially from in vitro and cell culture assays. 20 For example, a dose can be formulated in animal models to achieve a desired concentration or titer. Such information can be used to more accurately determine useful doses in humans.

25 Toxicity and therapeutic efficacy of the active ingredients described herein can be determined by standard pharmaceutical procedures in vitro, in cell cultures or experimental animals. The data obtained from these in vitro and cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage may vary depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. (See e.g., Fingl, 30 et al., 1975, in "The Pharmacological Basis of Therapeutics", Ch. 1 p.1).

Dosage amount and interval may be adjusted individually to brain or blood levels of the active ingredient are sufficient to induce or suppress the biological effect

(minimal effective concentration, MEC). The MEC will vary for each preparation, but can be estimated from in vitro data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. Detection assays can be used to determine plasma concentrations.

5 Depending on the severity and responsiveness of the condition to be treated, dosing can be of a single or a plurality of administrations, with course of treatment lasting from several days to several weeks or until cure is effected or diminution of the disease state is achieved.

10 The amount of a composition to be administered will, of course, be dependent on the subject being treated, the severity of the affliction, the manner of administration, the judgment of the prescribing physician, etc.

15 Compositions of the present invention may, if desired, be presented in a pack or dispenser device, such as an FDA approved kit, which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. The pack or dispenser may also be accommodated by a notice associated with the container in a form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals, which notice is reflective of approval by the agency of the form of the compositions or human 20 or veterinary administration. Such notice, for example, may be of labeling approved by the U.S. Food and Drug Administration for prescription drugs or of an approved product insert. Compositions comprising a preparation of the invention formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of an indicated condition, as is further detailed 25 above.

As used herein the term "about" refers to $\pm 10\%$.

30 As used herein the term "method" refers to manners, means, techniques and procedures for accomplishing a given task including, but not limited to, those manners, means, techniques and procedures either known to, or readily developed from known manners, means, techniques and procedures by practitioners of the chemical, pharmacological, biological, biochemical and medical arts.

As used herein, the term "treating" includes abrogating, substantially inhibiting, slowing or reversing the progression of a condition, substantially ameliorating clinical or aesthetical symptoms of a condition or substantially preventing the appearance of clinical or aesthetical symptoms of a condition.

5 It is appreciated that certain features of the invention, which are, for clarity, described in the context of separate embodiments, may also be provided in combination in a single embodiment. Conversely, various features of the invention, which are, for brevity, described in the context of a single embodiment, may also be provided separately or in any suitable subcombination or as suitable in any other described 10 embodiment of the invention. Certain features described in the context of various embodiments are not to be considered essential features of those embodiments, unless the embodiment is inoperative without those elements.

15 Various embodiments and aspects of the present invention as delineated hereinabove and as claimed in the claims section below find experimental support in the following examples.

EXAMPLES

Reference is now made to the following examples, which together with the above descriptions illustrate some embodiments of the invention in a non limiting fashion.

20 Generally, the nomenclature used herein and the laboratory procedures utilized in the present invention include molecular, biochemical, microbiological and recombinant DNA techniques. Such techniques are thoroughly explained in the literature. See, for example, "Molecular Cloning: A laboratory Manual" Sambrook et al., (1989); "Current Protocols in Molecular Biology" Volumes I-III Ausubel, R. M., ed. 25 (1994); Ausubel et al., "Current Protocols in Molecular Biology", John Wiley and Sons, Baltimore, Maryland (1989); Perbal, "A Practical Guide to Molecular Cloning", John Wiley & Sons, New York (1988); Watson et al., "Recombinant DNA", Scientific American Books, New York; Birren et al. (eds) "Genome Analysis: A Laboratory Manual Series", Vols. 1-4, Cold Spring Harbor Laboratory Press, New York (1998); 30 methodologies as set forth in U.S. Pat. Nos. 4,666,828; 4,683,202; 4,801,531; 5,192,659 and 5,272,057; "Cell Biology: A Laboratory Handbook", Volumes I-III Cellis, J. E., ed. (1994); "Culture of Animal Cells - A Manual of Basic Technique" by Freshney, Wiley-

Liss, N. Y. (1994), Third Edition; "Current Protocols in Immunology" Volumes I-III Coligan J. E., ed. (1994); Stites et al. (eds), "Basic and Clinical Immunology" (8th Edition), Appleton & Lange, Norwalk, CT (1994); Mishell and Shiigi (eds), "Selected Methods in Cellular Immunology", W. H. Freeman and Co., New York (1980);
5 available immunoassays are extensively described in the patent and scientific literature, see, for example, U.S. Pat. Nos. 3,791,932; 3,839,153; 3,850,752; 3,850,578; 3,853,987; 3,867,517; 3,879,262; 3,901,654; 3,935,074; 3,984,533; 3,996,345; 4,034,074; 4,098,876; 4,879,219; 5,011,771 and 5,281,521; "Oligonucleotide Synthesis" Gait, M. J., ed. (1984); "Nucleic Acid Hybridization" Hames, B. D., and Higgins S. J., eds. (1985); "Transcription and Translation" Hames, B. D., and Higgins S. J., eds. (1984); "Animal Cell Culture" Freshney, R. I., ed. (1986); "Immobilized Cells and Enzymes" IRL Press, (1986); "A Practical Guide to Molecular Cloning" Perbal, B., (1984) and "Methods in Enzymology" Vol. 1-317, Academic Press; "PCR Protocols: A Guide To Methods And Applications", Academic Press, San Diego, CA
10 (1990); Marshak et al., "Strategies for Protein Purification and Characterization - A Laboratory Course Manual" CSHL Press (1996); all of which are incorporated by reference as if fully set forth herein. Other general references are provided throughout this document. The procedures therein are believed to be well known in the art and are provided for the convenience of the reader. All the information contained therein is
15 incorporated herein by reference.
20

EXAMPLE 1

Identification of active peptides from exons 6 and 10 of CD44

Materials and Methods

25 All peptides were synthesized by LifeTein (South Plainfield, NJ, USA) at >95% purity. N2A mouse neuroblastoma and SK-N-SH human neuroblastoma (ATCC) were maintained in Dulbecco's modified Eagle's medium supplemented with 10 % fetal calf serum, L-Glut and 1 % Penicillin-Streptomycin (Beit Haemek, Israel). Cells were maintained in an incubator at 37 °C with 5 % CO₂. SK-N-SH cells were treated with 3
30 μM RA (Sigma-Aldrich) for 5 days prior to each experiment to allow the cells to differentiate towards neuronal cells. Cells were grown in 24 wells plate and treated with Aβ peptides for 48 hrs or MPTP (Sigma-Aldrich, 24 hrs) afterwhich they were

subjected to to the XTT viability assay. The XTT viability assay is based on the ability of metabolic active cells to reduce the tetrazolium salt XTT to orange colored compounds of formazan. The intensity of the water soluble dye is proportional to the number of metabolic active cells. XTT (Beit Haemek, Israel) was added to the cells 5 following treatment after 1:3 dilution with growth medium and incubated for 1-2 hours. Absorbance was measured at 420 nm. The viability assay was followed by caspase 3 assay (EnzChek Caspase3 Assay kit, Invitrogen, Carlsbad, Ca, USA) according to the manufacturer's instructions

Results

10 In vitro and in vivo loss of function experiments using siRNA, indicate that CD44V6 and CD44V10 are playing a role in neurodegenerative disorders such as AD, PD and ALS (unpublished data). CD44S and splice variant isoforms were shown to participate in multiple protein-protein interactions, including in signal transduction pathways (reviewed by Ponta et al [9]). Therefore the present inventors envisioned that 15 such interactions may mediate CD44V6 and CD44V10 function in neuronal cell death and that peptides derived from V6 or V10 exons sequences may serve as an agent for disruption of these interactions. Indeed it was shown that a small pentapeptide (NRWHE- SEQ ID NO: 1) derived from the human V6 sequence is capable of inhibiting hepatocyte growth factor (HGF) signal transduction through Met tyrosine 20 kinase receptor [24].

In order to identify conserved sequences that may help to identify such peptides, the present inventors have made a multiple species sequence alignment for both V6 and V10 sequences (Figure 1). Based on the conserved sequences, synthesized several peptides that cover different regions in V6 and V10 exons were synthesized (Figure 2). 25 The peptides were tested for their effect on cell death of neuroblastoma cell lines N2A (mouse) and SK-N-SH (human). As an example shown in Figure 3, the effect of two peptides derived from mouse V6 sequence and two peptides derived from the human V10 sequence was tested, at three concentrations, on cell death of SK-N-SH cells induced by A β 1-42 peptide. Cellular viability was measured by reduction of XTT. 30 Shown are the relative viability percentage as 100% defined as the viability of the none-treated cells. All 4 peptides were found to confer at least partial protection to the cells compared to no peptide control (Figure 3). Surprisingly, the protection conferred the V6

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peptides was maximal at a concentration as low as 20 pM and the degree of protection was reduced at higher concentrations. Additional peptides were further synthesized (6-21 amino acids, Figure 2) derived from mouse V6 and human V10 sequences and their protection activity from A β 25-35 toxicity in N2A cells was compared (Figures 4A-B).

5 A β 25-35 mimics the toxicological and aggregational properties of the full-length peptide with increased potency [25]. At 1 nM concentration, most of the peptides protected the cells, at least partially, as measured by both viability assay (XTT, Figure 4A) and caspase 3 activity (as marker for apoptosis induction, Figure 4B). These peptides were also tested for protection from cell death induced by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP), a PD modeling neurotoxin which is metabolized to the toxic cation 1-methyl-4-phenylpyridinium (MPP $^+$). Indeed some of the peptides were found also to have a protective effect against MPTP at 1 μ M compared to control cells (without peptide). Results are presented in Figures 5A-B.

15

EXAMPLE 2

Structural/functional analyses of CD44 peptides

Materials and methods

All methods are the same as in Example 1 above. Peptides and 6-hydroxydopamine were purchased from Sigma-Aldrich (St. Lewis, USA).

Results

These finding prompted the present inventors to expand the peptide screen to smaller peptides derived from conserved regions in V6B, V10A and V10B human sequences (Table 1).

Table 1 (for Figures 6 and 7)

Name	Sequence	SEQ ID NO:
V6B1	TPKEDSH	16
V6B1_C4	EDSH	17
V10A/B_L	PVTSAKTGSFGVTAVTV	18
V10A/B_S	PVTSAKTGSFG	19
V10A2_C7	PVTSAKT	20
V10A2_C10	TFIPVTSAKT	21
V10A1_N12	TTLLEGYTSHYP	22
V10A1_N8	TTLLEGYT	23
V10A1_N6	TTLLEG	24
V10A1_N+2_6	LLEGYT	25

Name	Sequence	SEQ ID NO:
V10A1_N+4 (also referred to as P26)	EGYTSHPY	26
V10A1_EGYT	EGYT	27
V10B3_N7	SLSGDQDT	28
V10B3_N6	SLSGDQD	29
V10B3_N5	SLSGD	30
V10B2.5_6	NVNRSL	31
V10B2.5_9	NVNRSLSGD	32
V10B2_N10	DSNSNVNRSL	33
V10B1_8 (also referred to as P34)	FGVTAVTV	34
V10B1_7	FGVTAVT	35

These peptides were screened for their protective effect on 6-hydroxydopamine (6-OHDA) treated N2A cells at 1 pM concentration. 6-OHDA is a dopamine analogue that is commonly used in model systems to mimic Parkinson's disease in vitro and in vivo. 6-OHDA induces apoptosis through release of reactive oxygen species (ROS) and by a possible direct effect on the mitochondrial respiratory chain. The results show differential effect of the peptides on cellular viability and caspase 3 activation on 6-OHDA treated cells (Figures 6A-B). The protective effect of some of the peptides was tested in a wide dose range. For example, for the 8 mer peptide V10A1_N+4, the optimal protective dose was found to be around the fM concentration (Figure 6C). The peptide panel was also tested for their effect on A β (25-35) induced toxicity in SK-N-SH human neuroblastoma cells at pM concentration. The profile of peptides effective against the A β stress was strikingly different than the 6-OHDA-effective peptides (Figures 7A-B).

15

EXAMPLE 3

In vivo effect of some peptides of the invention in an in vivo Parkinson's model

Materials and Methods

20 C57BL mice (all male, age 8-12 weeks, obtained from Harlan Laboratories Israel) were administered with 18 mg/kg MPTP (Sigma-Aldrich) at a dosage volume of

100 μ l/mouse by intraperitoneal (IP) injection twice daily, 3 hrs apart on days 1 and 2. On study day 0 and for 8 consecutive days, mice were administered intranasally with PBS (vehicle) or one of the peptides at 1 mg/kg in 12 μ l/mouse. All peptides used in in vivo studies were synthesized by LifeTein and were modified with N terminal acetylation and C terminal amidation. For intranasal instillation, each mouse was mildly anesthetized (2.5% Isoflurane) then restrained and held with the neck parallel to the table while a total volume of 12 μ l was administered into the nostril. Six (6) μ l was administered to the left nostril as two 3 μ l drops, followed by a 15 sec hold, and 6 μ l was administered to the right nostril as two 3 μ l drops, followed by a 15 seconds hold. On day 8 the animals were euthanized by cervical dislocation. Immediately after euthanasia, the brains were removed and the striata dissected (left and right striatum were pooled), weighed and frozen in dry ice. The striata samples were homogenized in a solution containing 0.1 M perchloric acid and 10 ng/ml 3,4 dihydroxybenzylamine (DHBA) by 5 seconds sonication at 80 W. The supernatants of each tissue extract were injected directly to HPLC pump (Jasco PU- 2080Plus) onto a reverse phase column (GL-Science, Inertsil ODS-2 5um 4.6x150mm at room temperature) coupled to an electrochemical detector Coulochem II ESA with a conditioning cell model 5021 and analytical cell model 5011. The working potential was set to 0.35V on the conditioning cell and 0.1V and - 0.35V on the analytical cell. The mobile phase was 0.05M monobasic sodium phosphate ,with 80mg/L EDTA, 125 mg/L heptane sulfonic acid, 55ml of methanol and 50ml of acetonitrile pH=2.7. The flow rate was 1.5 ml/min. The dopamine, DOPAC and HVA values were normalized to the lysate protein concentration (BCA kit, Pierce).

25

Results

The most effective peptides against MPTP and 6-OHDA were selected for in vivo PD model namely MPTP injection in mice. In this model, repeated intraperitoneal injections of MPTP (18 mg/kg at 3-h intervals on two consecutive days) result in dopaminergic neuronal death in the substantia nigra and dopamine depletion of the striatum. The peptides (or vehicle) were applied twice daily by intranasal instillation at 1 mg/kg starting 1 day prior to exposure to MPTP and continued until the end of the study. 7 days after the injection of MPTP mice were sacrificed and striatal levels of dopamine (DA), 3,2-dihydroxyphenylacetic acid (DOPAC) and homovanillic acid

(HVA) were evaluated by HPLC. 9 peptides were tested in 8 groups and the results of DA, DOPAC and HVA levels are shown in Table 2.

Specifically Table 2 illustrates the effect of human V6 and V10-derived peptides (1 mg/kg, intranasal administration) on the striatal level of dopamine and metabolites in 5 MPTP-treated mice. Mice were treated according to the protocol described in the Materials and Methods. Striatal levels of DA, DOPAC and HVA were determined by HPLC, divided by the total protein content value and normalized to the levels obtained in vehicle treated mice. Groups 1 and 3 treated with 1:1 mix of normal designated and retro-inverso analog (D amino acids at reversed sequence). Group 6 was treated with 10 1:1 mix of 10B2.5_6 and 10B2.5_9 peptides.

Table 2

	SEQ ID NO	Name of peptide	DA	DOPAC	HVA
1	17	hV6B1_C4	20.7	41.7	16.6
2	19	V10A/B_S	26.6	27.2	13.7
3	24	V10A1_N6	17.7	24.3	-0.7
4	26	V10A1_N+4	58.0 *	61.8 *	28.5
5	15	hV10B3	18.1	22.5	44.2 *
6	31 and 32	10B2.5_6+10B2.5_9	13.6	35.1	53.0 *
7	34	10B1_8	46.6 *	75.9 **	83.3 **
8	12	hV10A2	24.7	54.8 *	73.4 **

* p value <0.05

** p value <0.01

15 Treatment with two 8-mer, V10-derived peptides namely V10A1_N+4 (SEQ ID NO: 26) and 10B1_8 (SEQ ID NO: 34) had a significantly positive effect on striatal DA level compared to the vehicle control. These peptides, as well as three other peptides (hV10B3 – SEQ ID NO: 15; V10A2 – SEQ ID NO:12; 10B2.5 _6 – SEQ ID NO: 31; and 10B2.5_9 – SEQ ID NO: 32), also increased significantly the levels of at least one 20 of the 2 DA metabolites, DOPAC and HVA (Table 2, above). These results indicate that peptides derived from CD44 V10 are able to protect dopaminergic neurons from MPTP in vivo and suggest that these peptides be developed as novel drugs for Parkinson's disease.

EXAMPLE 4**Identification of active subsequences within the peptides of some embodiments of the invention****5 Results**

V10A1_N+4 (hereunder "P26", SEQ ID NO: 26) was chosen for further structure-function analysis. Peptides which were derived from this peptide were synthesized and shown on Table 3, below.

Table 3

Name	SEQ ID NO:	Sequence
26-1	36	EGYTS HY
26-2	37	EGYTS H
26-3	38	EGYTS
26-4	39	GYTSHYP
26-5	40	YTS HYP
26-6	41	TSHYP
26-7	42	GYTSHY
26-8	43	QGYTSHYP
26-9	44	EGYTSAYP
26-RI	45	*P*Y*H*S*T*Y*G*E
26-R	46	PYHSTYGE

10

* D amino acid

The peptides shown on Table 3 were tested for their effect in protecting N2A from 6-OHDA (Figure 8). This data indicates that some modifications in the P26 sequence can be tolerated while maintaining at least partial activity. Such modifications 15 include C-terminus truncation (26-1, 26-2 and 26-3), replacement of the N-terminal glutamic acid residue with glutamine and histidine at position 7 with alanine (26-8 and 26-9, respectively).

Most interestingly, a retro-inverso (RI, in which the primary sequence is reversed and D- rather than L- amino acids are used) P26 derivative (26-RI, SEQ ID NO: 26) also retain its neuroprotection activity. It is postulated that a retro-inverso peptides assume a side chain topology, in its extended conformation, similar to that of its native L-sequence and retaining the biological activities of the parent molecule while fully resistant to proteolytic degradation [Chorev, M. and M. Goodman, *Recent developments in retro peptides and proteins--an ongoing topochemical exploration*. Trends Biotechnol, 1995. 13(10): p. 438-45]. Therefore, it is contemplated that a minimal sequence for mediating neuroprotection is set forth in SEQ ID NO: 47 STYG-X (where X is E or Q) or retro configuration of same, whereby every amino acid can be D or L.

EXAMPLE 5

Intrahippocampal (IH)/intracerebroventricular (ICV) injection of V10A1_N+4

15 **(P26) and P34 peptides protects rats from A β ₍₁₋₄₂₎ damage**

Materials and Methods

Animals - Adult male Sprague Dawley rats were obtained from the Laboratory Animal Center of University of South China, Hengyang, Hunan, China. After arrival, the rats were housed individually in a temperature- and humidity-controlled environment with *ad libitum* access to food and water. Animals were maintained on a 20 12 hr light/dark schedule, with lights on at 7 A.M. After being housed, the rats were handled (5–6 min per rat per day) for 1 week to habituate them to the experimenter. Experiments were conducted according to the *National Institutes of Health Guide for the Care and Use of Laboratory Animals*, and experimental protocols were approved by 25 the University Animal Care and Use Committee.

Drugs - A β ₍₁₋₄₂₎ was purchased from Sigma-Aldrich (USA). Peptides obtained from LifeTein (USA).

Establishment of the Alzheimer disease (AD) model - A β ₍₁₋₄₂₎ (Sigma-Aldrich, USA), was dissolved in filtered PBS at the concentration of 6 μ g/ μ l, and the 30 solution kept at 37°C for 2 days before use. One microliter of the solution was injected by means of a microsyringe into the right hippocampus under sodium pentobarbital (55 mg/kg i.p.) anesthesia at the stereotaxic coordinates: AP= 3.6, ML=2.0, from Bregma

and DV=3.0 from the skull. Control rats were injected with 1 μ l of PBS. In the IH/ICV study, 1 μ l of the peptide or PBS was also injected into the right hippocampus at the same stereotaxic coordinates.

5 **Peptide administration** - For ICV injection, a microinjection cannula was planted into the lateral cerebral ventricle according to the following stereotaxic coordinates: AP= 1, ML=1.6, DV=3.8. Twenty-four hours after A β injection, rats received the first ICV injection of PBS or peptides at the designated doses. In the subcutaneous (SC) study, daily SC injections of PBS or peptide solution in PBS was done at 1 ml/rat to a dose of 1 mg/kg and lasted for 21 days.

10 **Novel object recognition task** - The NOR task was tested 21 days after A β injection.

The training apparatus was a black Plexiglas box (50×50×40 cm) placed in a sound-attenuating cabinet which was located in a brightly lit and isolated room. Illumination was provided by a 15 W white house light mounted on the ceiling of 15 cabinet, and a 65 dB background noise was supplied by a ventilation fan in the cabinet. The floor of the box was covered with sawdust. The objects used in the task were made of water-repellant materials such as glass and plastic with differences in shape and color. The sizes of the objects were about 6×6×8 cm. Two objects were always located in the back corners of the box. The location and objects were counterbalanced to control 20 for any preferences that the rats might have had for one of the corners or of the objects. The behavioral procedure involved two phases: training and retention test. During the training trial, the rat was placed in the box and allowed to explore two identical objects for 10 min and the total time spent exploring both objects was recorded. Exploration of an object was defined as pointing the nose to the object at a distance of <1 cm and/or 25 touching it with the nose. The sawdust was stirred and the box and the objects were cleaned with 40% ethanol solution between trials. Twenty-four hours after training trial (retention test trial), one copy of the familiar object and a new object were placed in the same location as stimuli during the training phase. The rat was placed in the box for 3 min and the time spent exploring each object and the total time spent exploring both 30 objects were recorded. The discrimination index used to assess memory was calculated as the difference in time exploring the novel and familiar object, expressed as the ratio of the total time spent exploring both objects.

Morris water maze test - consisted of a circular water tank (200 cm diameter, 60 cm height) filled with water (25 ± 1 °C) to a depth of 40 cm. Four equally spaced locations around the edge of the pool were used as start points, which divided the pool into 4 quadrants. An escape platform (10 cm in diameter) was placed in the pool 2 cm below the surface of water. The escape platform was placed in the middle of one of the randomly selected quadrants of the pool and kept in the same position throughout the entire experiment. Before the training started, the rats were allowed to swim freely into the pool for 120 s without platform. Animals received a training session consisting of 4 trials per session (once from each starting point) for 4 days, each trial having a ceiling time of 120 s and a trial interval of approximately 30 s. After climbing onto the hidden platform, the animals remained there for 30 s before commencement of the next trial. If the rat failed to locate the hidden platform within the maximum time of 120 s, it was gently placed on the platform and allowed to remain there for the same interval of time. The time taken to locate the hidden platform (latency in seconds) was measured. Twenty four hours after the acquisition phase, a probe test was conducted by removing the platform. Rats were allowed to swim freely in the pool for 120 s and the time spent in target quadrant, which had previously contained the hidden platform, was recorded. The time spent in the target quadrant indicated the degree of memory consolidation which had taken place after learning. The Morris water maze test was started 24 days after A β injection.

Statistical Analyses - Statistical analyses were performed using one-way ANOVA. Post-hoc comparisons were performed with the Fisher LSD Test (SigmaStat 3.2). All data were represented as mean \pm SEM. Significant level was set at $p<0.05$.

Results

In order to test the efficacy P26 and V10B1_8 (hereinunder P34) peptides in Alzheimer's disease (AD) model *in vivo*, these peptides were tested in A β ₍₁₋₄₂₎ microinjection rat model [Soto, C., et al., *Beta-sheet breaker peptides inhibit fibrillogenesis in a rat brain model of amyloidosis: implications for Alzheimer's therapy*. Nat Med, 1998. 4(7): p. 822-6]. In this model, rats received a single microinjection of 6 μ g A β ₍₁₋₄₂₎ into the right hippocampus. At the same time, 1 μ l of peptide solution or PBS (vehicle) was also injected into the same location. This treatment was followed by daily ICV injection of the peptides at different doses or

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vehicle (PBS). 21 days after A β injection the rats were tested for the novel object recognition assay (NOR, Figure 9). The results show that P26 at 100 ng/rat and P34 at 10 and 100 ng/rat increased significantly the discrimination index ($p<0.01$). The data indicates that ICV/IH administration of the P26 and P34 peptides prevent the toxic 5 effect of A β in vivo.

EXAMPLE 6

Peripheral injection of V10A1 N+4 (P26) peptide protects rats from A β ₍₁₋₄₂₎ damage

10 Material and Methods

Experimental procedures and subcutaneous injection (SC) is described in Example 5 above.

Results

15 Parenteral administration of the peptides by subcutaneous (SC) injection was assayed for protecting the rats from A β toxicity. Therefore the A β ₍₁₋₄₂₎ microinjection rat model was applied followed by daily SC injection of 1 mg/kg P26 peptide, as well as the control peptide (cont1: AVAVEAAG SEQ ID NO: 48, n=10-11). The 21 days injection period was followed by Morris water maze (MWM) and NOR memory assays. The results show that of the 3 SC-injected peptide only P26 improved significantly the 20 behavior in both assays (Figures 10 and 11). These results demonstrate that the neuroprotection effect of the P26 peptide is seen also when the peptide is given by parenteral administration, suggesting that the peptide is able to cross the blood-brain barrier and reach sufficient concentration in the relevant brain regions.

25

EXAMPLE 7

pharmacokinetics of P26 and P26-IR

Materials and Methods

30 The pharmacokinetics of the peptides was evaluated following SC peptide administration in male Sprague Dawley rats. Peptide solution was prepared using phosphate buffer saline (pH 7.4) as vehicle and was administered through subcutaneously at the dose of 1 mg/kg with dosing volume of 2 mL/kg.

Blood samples were collected at 0 (pre-dose) and 0.17, 0.5, 1, 2, 4, 8 and 24 hours following administration. At each time point, approximately 0.25 mL of blood was withdrawn through jugular vein of the cannulated rats and transferred to a pre-labeled microfuge tube containing 200 mM K₂EDTA (20 µL per mL of blood). Following sampling equal volume of heparinized saline was flushed in to catheter. Blood samples were centrifuged at 5000g for 5 minutes at 4 ± 2°C. All the plasma samples were stored below -70 °C until analysis. A fit-for-purpose LC-MS/MS method was used for the quantification of the peptides in plasma samples. The lower limit of quantification (LLOQ) was 22.34 ng/mL.

Results

The pharmacokinetics of P26 and P26-IR were tested in rats by subcutaneous injection at 1 mg/kg (Figure 12). P26-RI demonstrated improved SC pharmacokinetics with an apparent C_{max} of 1227 ng/ml compared to 130 ng/ml for P12. This 10 fold improvement by P26-IR analogue, taken together with its activity in vitro, suggests that P26-IR can be used at lower doses compared to P26.

Although the invention has been described in conjunction with specific embodiments thereof, it is evident that many alternatives, modifications and variations will be apparent to those skilled in the art. Accordingly, it is intended to embrace all such alternatives, modifications and variations that fall within the spirit and broad scope of the appended claims.

All publications, patents and patent applications mentioned in this specification are herein incorporated in their entirety by reference into the specification, to the same extent as if each individual publication, patent or patent application was specifically and individually indicated to be incorporated herein by reference. In addition, citation or identification of any reference in this application shall not be construed as an admission that such reference is available as prior art to the present invention. To the extent that section headings are used, they should not be construed as necessarily limiting.

The terms "comprises", "comprising", "includes", "including", "having" and their conjugates mean "including but not limited to".

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(other references are recited in the application)

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CLAIMS

1. An isolated peptide comprising:
 - (i) at least 5 amino acids of a CD44V10 amino acid sequence and no more than 15 amino acids of said CD44V10 amino acid sequence, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NOS: 49 or 50, the peptide comprising a neuroprotective activity; or
 - (ii) at least 3 amino acids of a CD44V6 amino acid sequence and no more than 20 amino acids of said CD44V6 amino acid sequence, the peptide comprising a neuroprotective activity, with the proviso that the peptide does not consist of the amino acid sequence as set forth in SEQ ID NO: 1, 51 or 52,
- and wherein said peptide of (i) comprises an amino acid sequence of formula I:

$X_1\text{-G\text{-}Y\text{-}T\text{-}S}$, wherein X_1 is any of a glutamic acid or glutamine;

and wherein said peptide of (ii) comprises a core sequence $X_1\text{-}X_2\text{-S\text{-}H}$, wherein X_1 and X_2 are acidic amino acids.
2. The isolated peptide of claim 1, wherein said peptide of (i) comprises an amino acid sequence as set forth in SEQ ID NOS: 12, 15, 19-26 28-45 or 46.
3. The isolated peptide of claim 1, wherein said peptide of (i) comprises an amino acid sequence as set forth in SEQ ID NOS: 12, 15, 26, 31, 32, 34, 45 and 46.
4. The isolated peptide of any one of claims 1 to 3, wherein said peptide of (ii) comprises an amino acid sequence as set forth in SEQ ID NOS: 2-7, 16 or 17.
5. The isolated peptide of claim 4, wherein said amino acid sequence comprises peptidomimetics.
6. The isolated peptide of claim 5, wherein said peptidomimetics comprises a retro-inverso mimetic.
7. The isolated peptide of any one of claims 1 to 6, wherein said peptide comprises said peptide of (i) and wherein said sequence is as set forth in SEQ ID NO: 26, 45 or 46.

8. The isolated peptide of any one of claims 1 to 7, wherein said CD44V10 amino acid sequence is a human CD44V10 amino acid sequence, and wherein said CD44V6 amino acid sequence is a human CD44V6 amino acid sequence.

9. The isolated peptide of claim 1, wherein X_1 of said peptide of (ii) comprises glutamic acid.

10. The isolated peptide of claim 4, wherein X_2 of said peptide of (ii) comprises aspartic acid.

11. The isolated peptide of any one of claims 1 to 10, wherein the peptide is attached to a cell penetrating agent.

12. The isolated peptide of any one of claims 1 to 11, wherein the peptide is no longer than 20 amino acids.

13. A pharmaceutical composition comprising as an active agent the isolated peptide of any one of claims 1 to 12 and a pharmaceutically effective carrier.

14. The isolated peptide of any one of claims 1 to 12 or the pharmaceutical composition of claim 13, formulated for subcutaneous or intranasal delivery.

15. A method of treating a neurodegenerative disorder in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of the isolated peptide of any one of claims 1 to 12 or claim 14 or the pharmaceutical composition of claim 13 or claim 14.

16. The method of claim 15, wherein said neurodegenerative disorder is selected from the group consisting of Parkinson's disease, Multiple Sclerosis, ALS, multi-system atrophy, Alzheimer's disease, stroke, traumatic brain injury, progressive supranuclear palsy, fronto-temporal dementia with parkinsonism linked to chromosome 17 and Pick's disease.

17. The method of claim 15, wherein said neurodegenerative disease is Parkinson's disease.

18. The method of claim 15, wherein the peptide comprises said peptide of (i) as defined in claim 1, and wherein said sequence is selected from the group consisting of SEQ ID NOs: 12, 15, 26, 31, 32 and 34; and wherein said neurodegenerative disorder is Parkinson's disease.

19. The method of claim 15, wherein said neurodegenerative disease is Alzheimer's disease.

Date: 14 November 2016

FIG. 1A

Multiple sequence alignment of CV44V6 exon

Human	P	S	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	G	N	R	W	H	E	G	Y	R	Q	T	P	K	S	T	T	G	T	A				
Chimp	P	S	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	G	N	R	W	H	E	G	Y	R	Q	T	P	R	E	D	S	H	S	T	T	G	T	A
Macaque	P	S	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	E	N	G	W	S	G	G	Y	P	Q	T	P	K	E	D	S	H	S	T	A	G	T	A
Wild boar	P	S	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	E	N	G	W	S	G	G	Y	P	Q	T	P	K	E	D	S	H	S	T	A	G	T	A
Dog	P	S	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	E	N	G	W	S	G	G	Y	P	Q	T	P	K	E	D	S	H	S	T	A	G	T	A
Rabbit	P	N	S	T	T	E	E	T	A	T	Q	K	E	Q	W	V	G	N	R	W	H	G	G	N	L	Q	Y	P	T	S	E	D	H	S	T	E	G	T	T
Rat	P	N	S	T	T	E	E	T	A	T	Q	K	E	Q	W	F	E	N	E	W	Q	G	K	N	P	P	T	P	S	E	D	H	V	T	E	G	T	T	
Mouse	P	N	S	T	A	E	A	T	A	T	Q	E	T	W	F	Q	N	G	W	Q	N	G	K	N	P	P	T	P	S	E	D	H	V	T	E	G	T	T	

FIG. 1B

Multiple sequence alignment of CD44V10 exon

Human	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Chimp	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Macaque	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Wild Boar	R	N	D	G	R	G	G	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G		
Rabbit	R	T	H	G	R	G	G	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G		
Hamster	R	S	G	G	K	D	G	R	G	G	R	D	P	N	H	S	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G
Rat	R	S	G	A	R	G	G	S	L	P	R	D	T	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G								
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V10	C6 Junction																																																																												
Human	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Chimp	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Macaque	R	N	D	V	I	G	G	R	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G	
Wild Boar	R	T	H	G	R	G	G	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G		
Rabbit	R	T	H	G	R	G	G	R	D	P	N	H	S	G	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G		
Hamster	R	S	G	G	K	D	G	R	G	G	R	D	P	N	H	S	T	T	L	E	G	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G
Rat	R	S	G	A	R	G	G	S	L	P	R	D	T	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G								
Mouse	R	S	G	A	R	D	R	G	G	S	L	P	T	T	S	H	Y	P	H	T	K	E	S	R	T	F	I	P	V	T	I	P	V	T	S	A	K	T	G	M	T	A	V	T	V	-	G	D	S	N	S	N	V	N	R	S	L	S	G	D	D	T	F	H	-	P	S	G	G								

FIG. 2

Mouse V6 peptides

V6 (SEQ ID NO: 2) PNSTAAATQQETWFGQNGKNNPPTPSEDSHVTEGTT
 V6A (SEQ ID NO: 3) PNSTAAATQQETWFGQNGW
 V6A1 (SEQ ID NO: 4) PNSTAAATQQETWFGQNGW
 V6A2 (SEQ ID NO: 5) WFQNGW) TPSEDSHVTEGTT
 V6B (SEQ ID NO: 6) TPSEDSH
 V6B1 (SEQ ID NO: 7) TPSEDSH

Human V10 peptides

V10 TTLLEGYTSHYPHPTKESRTFIPVTSAKTGSFGVITAVTVGDSNSNVNRSLSGDQDTFHPSGG
 (SEQ ID NO: 8)

V10A TTLLEGYTSHYPHPTKESRTFIPVTSAKT
 (SEQ ID NO: 9) GSFGVITAVTVGDSNSNVNRSLSGDQDTFHPSGG

V10B (SEQ ID NO: 10)

V10A1 TTLLEGYTSHYPHPTKESRTFIPVTSAKT
 (SEQ ID NO: 11) GSFGVITAVTVGDSNSNVNRSLSGDQDTFHPSGG

V10A2 (SEQ ID NO: 12) TKEESRTFIPVTSAKT

V10B1 (SEQ ID NO: 13) GSFGVITAVTVGDSNSNVN
 DSNSNVNRSLSGDQDTFHPSGG

V10B2 (SEQ ID NO: 14) DSNSNVNRSLSGDQDTFHPSGG

V10B3 (SEQ ID NO: 15) SLSGDQDTFHPSGG

FIG. 3

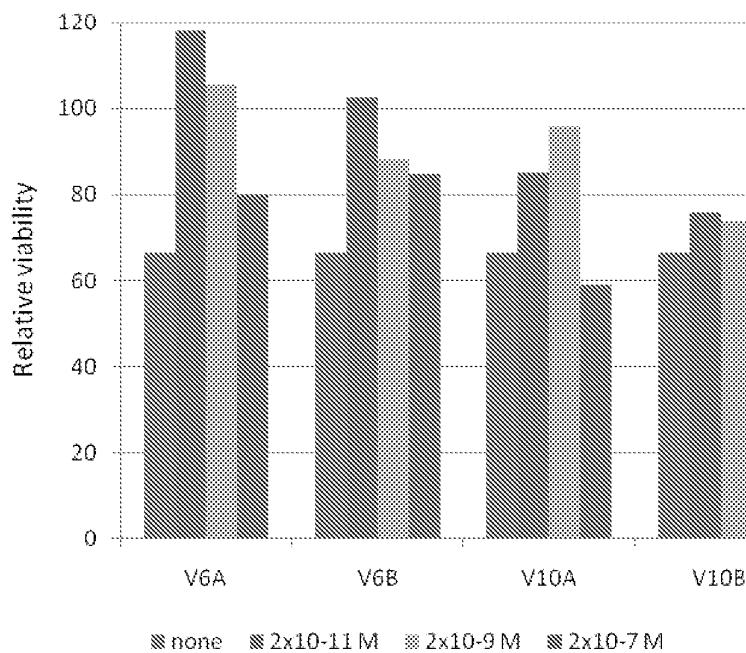


FIG. 4A

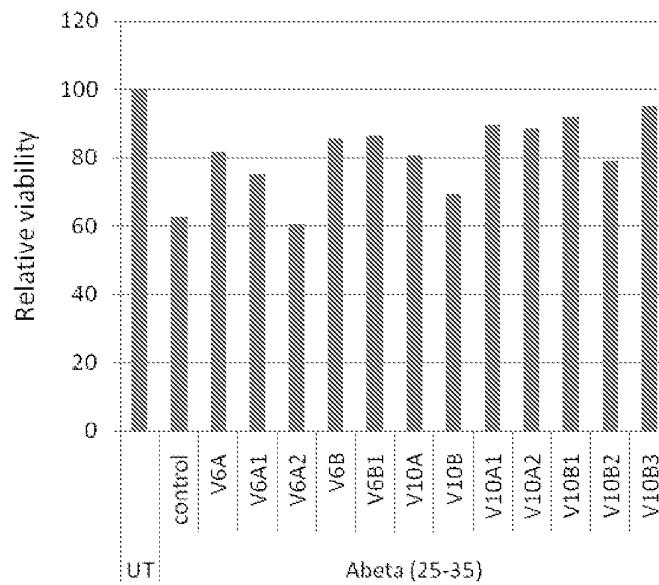


FIG. 4B

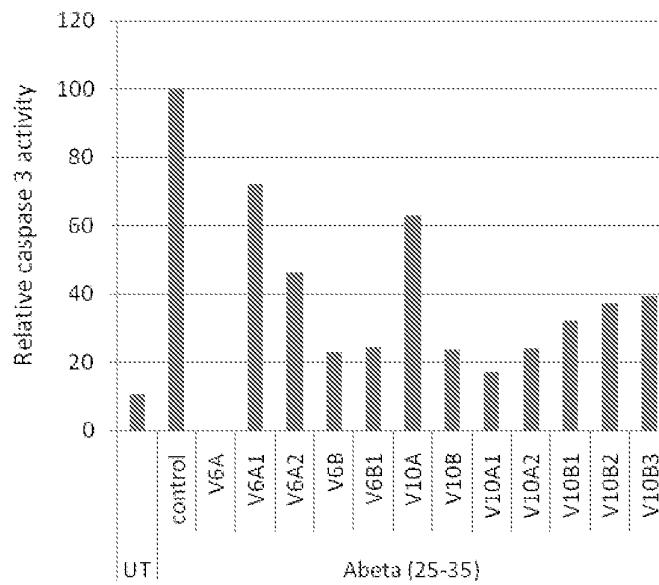


FIG. 5A

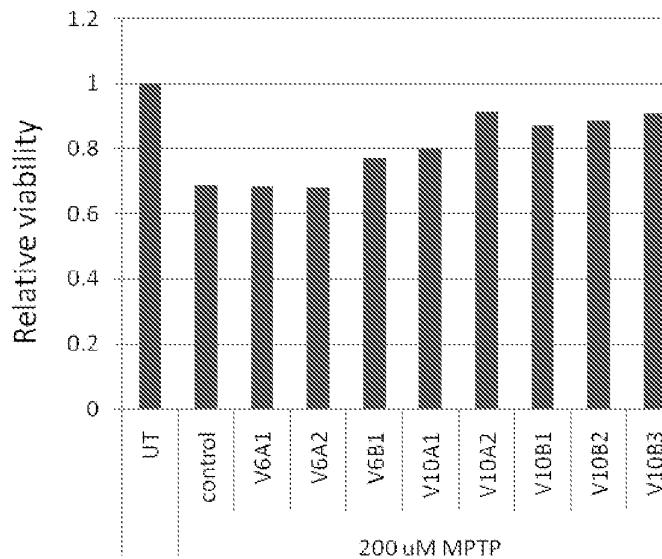


FIG. 5B

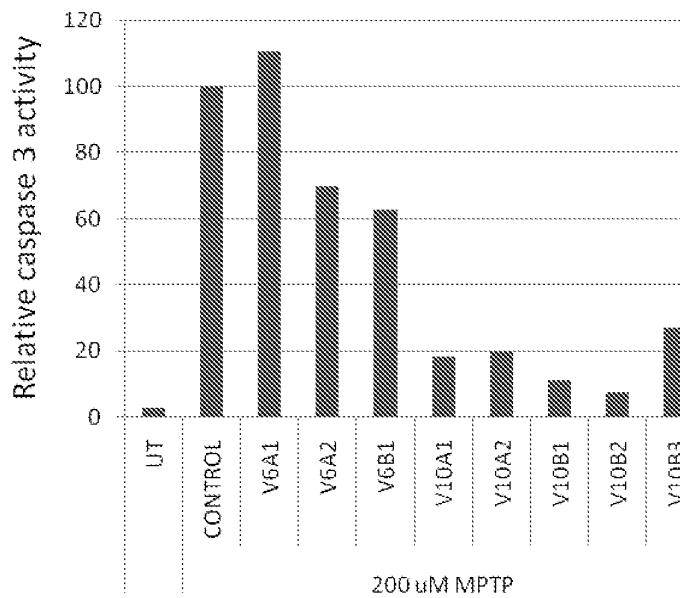


FIG. 6A

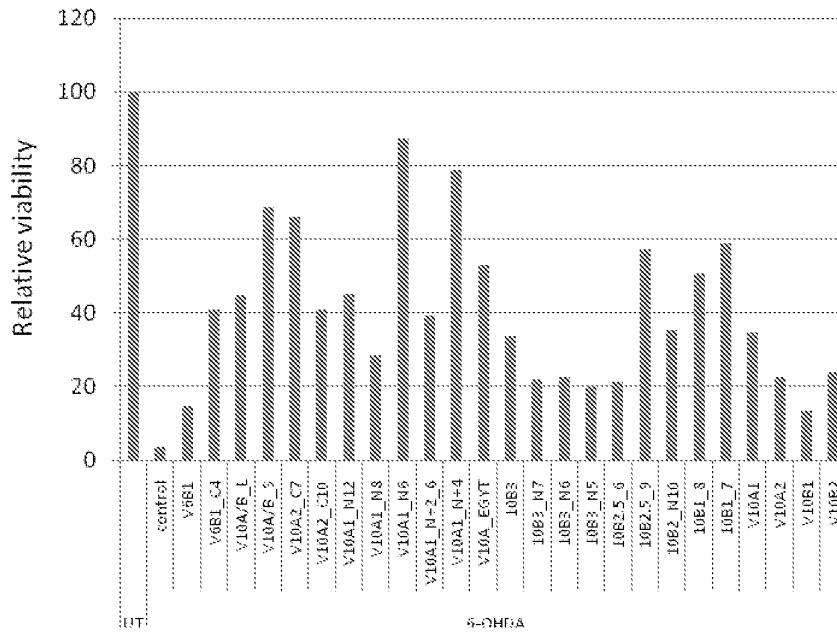


FIG. 6B

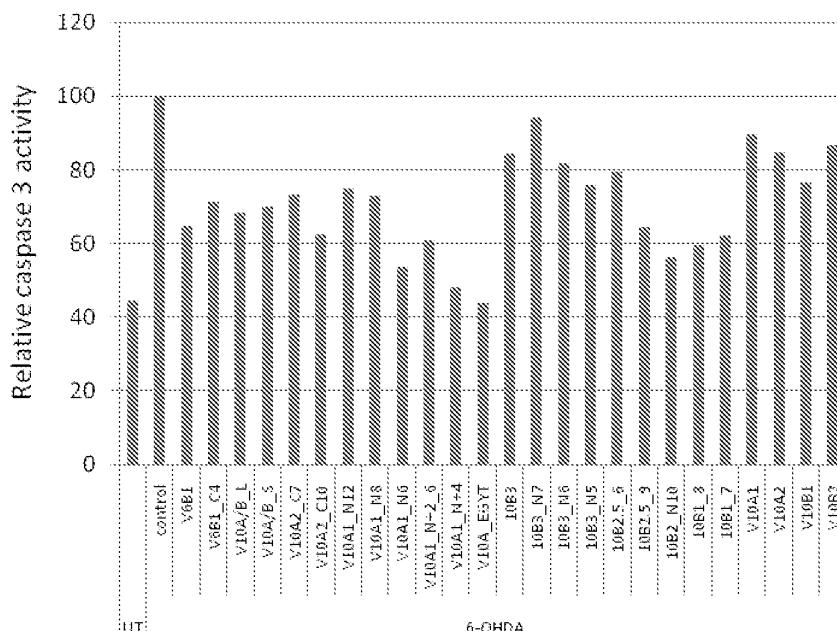


FIG. 6C

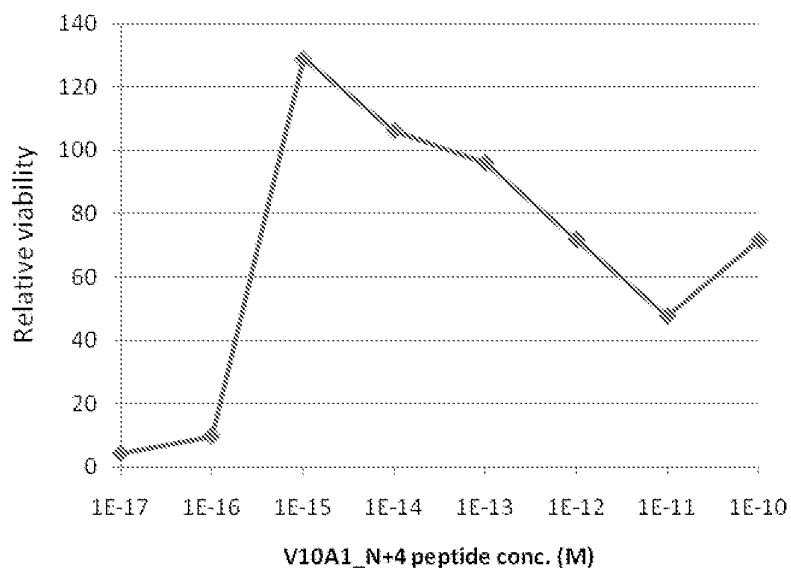


FIG. 7A

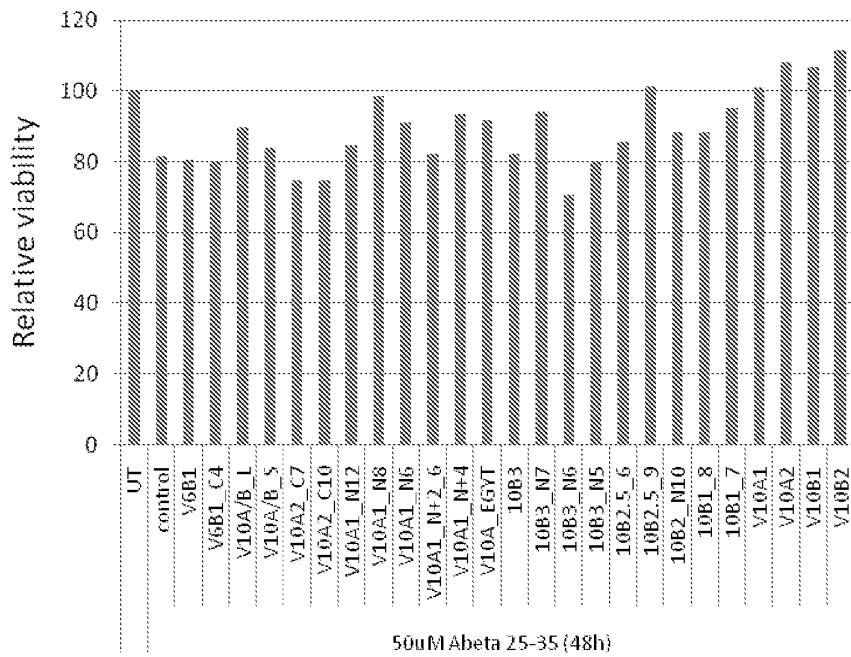


FIG. 7B

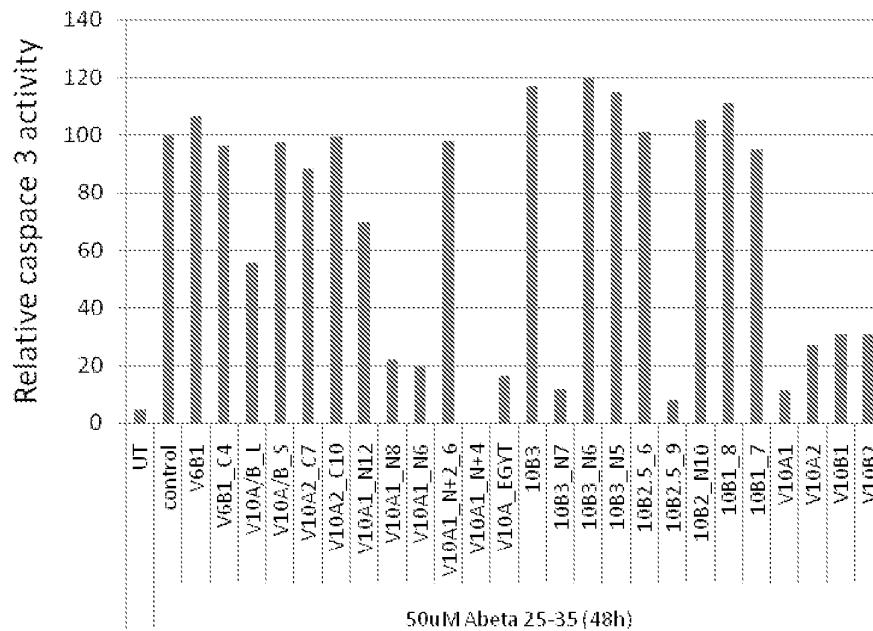


FIG. 8A

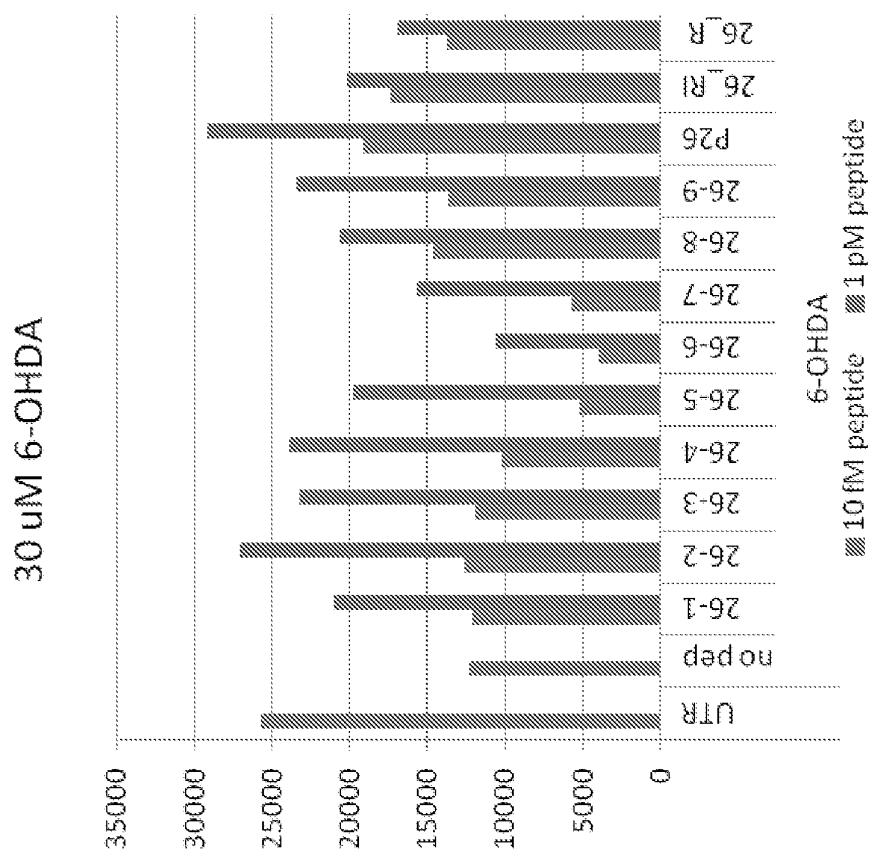


FIG. 8B

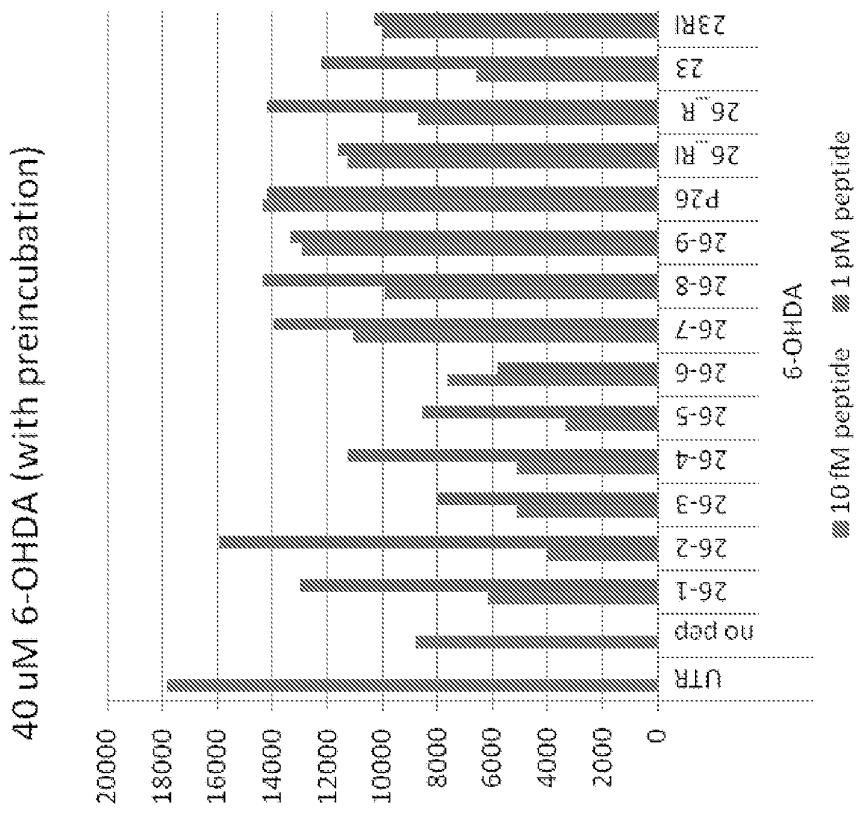


FIG. 9

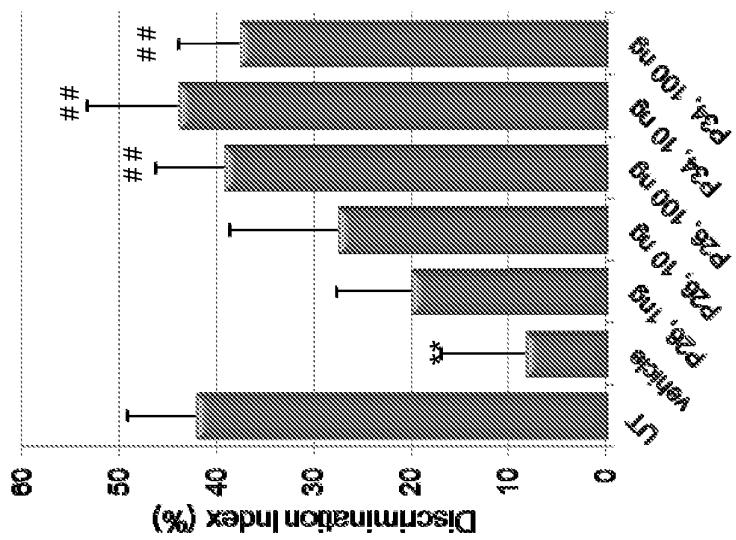
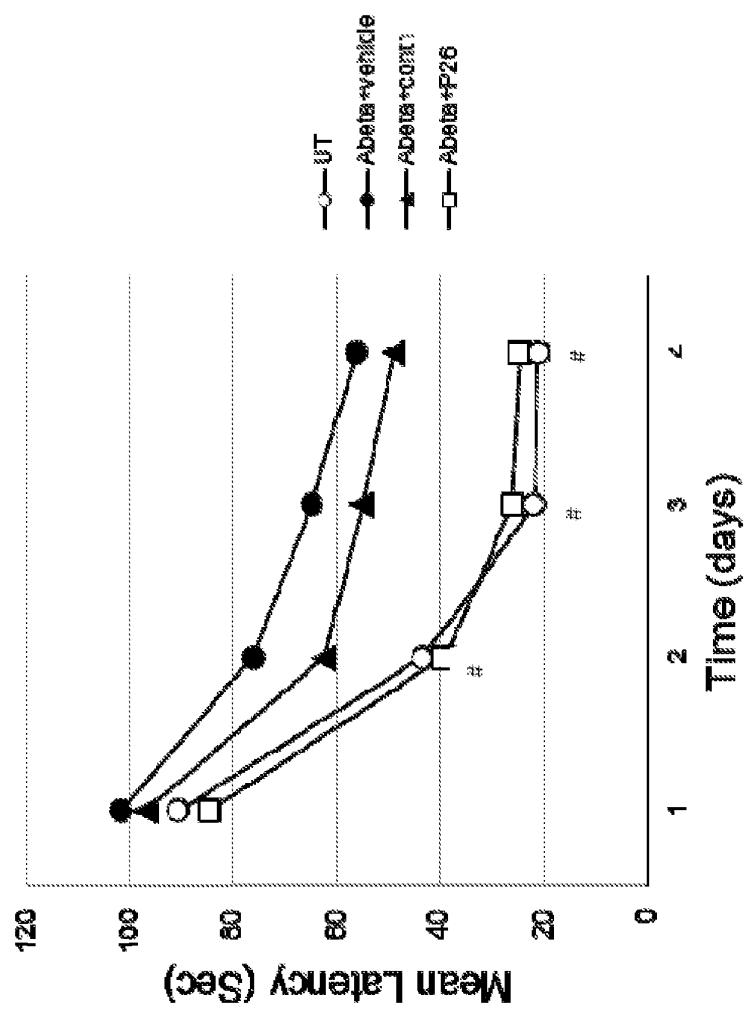


FIG. 10A



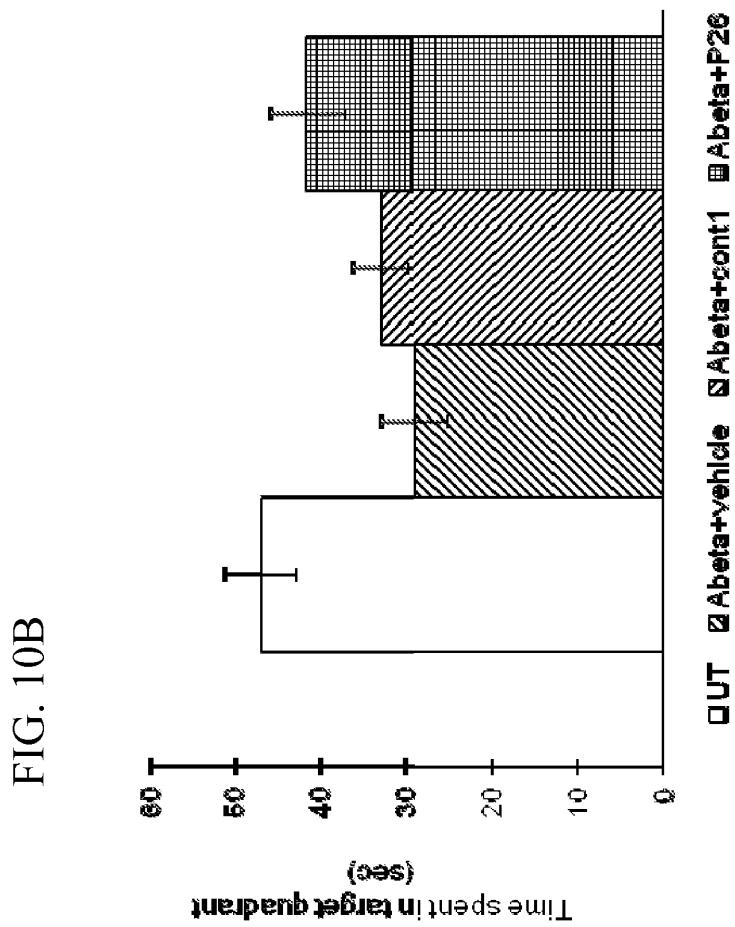


FIG. 11

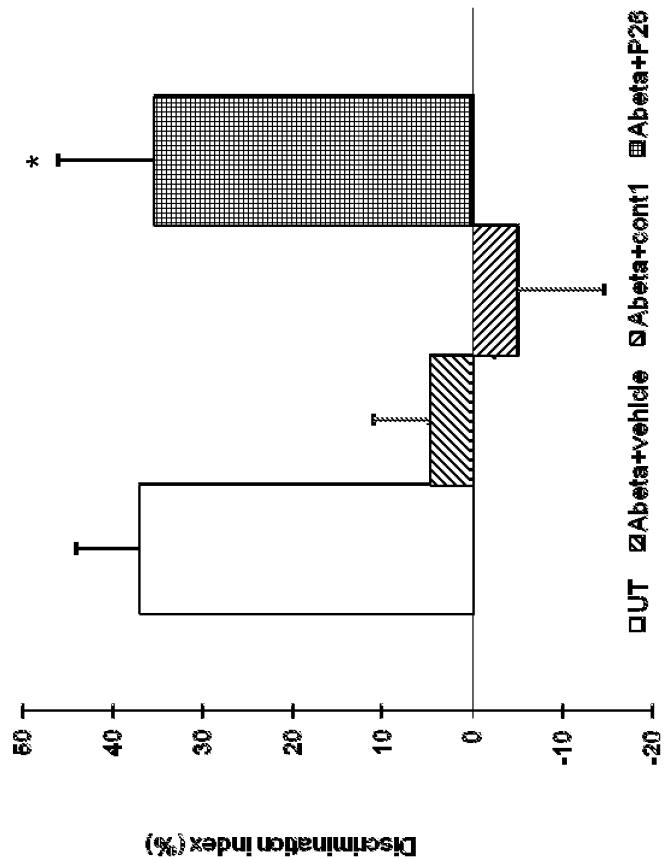
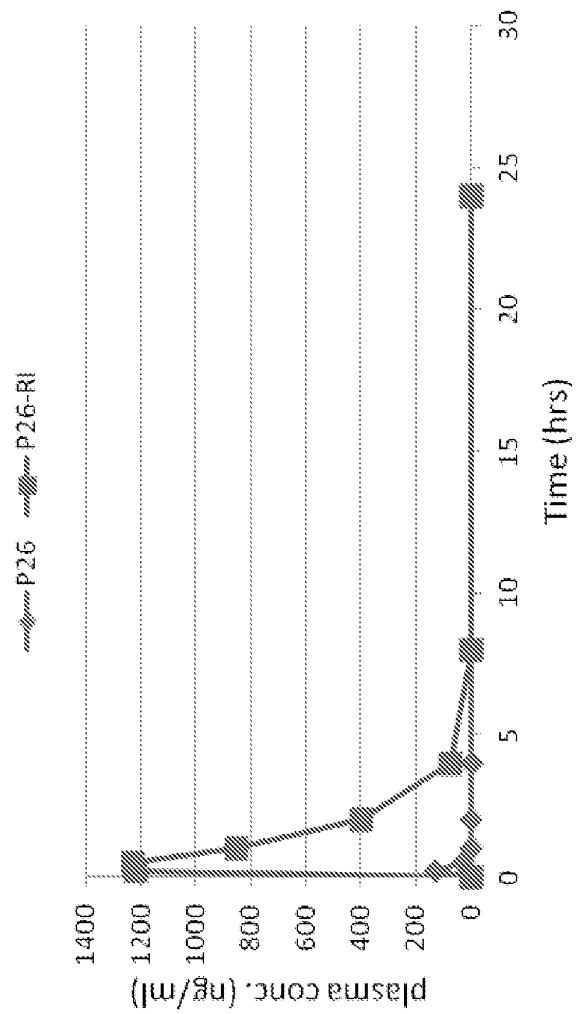


FIG. 12



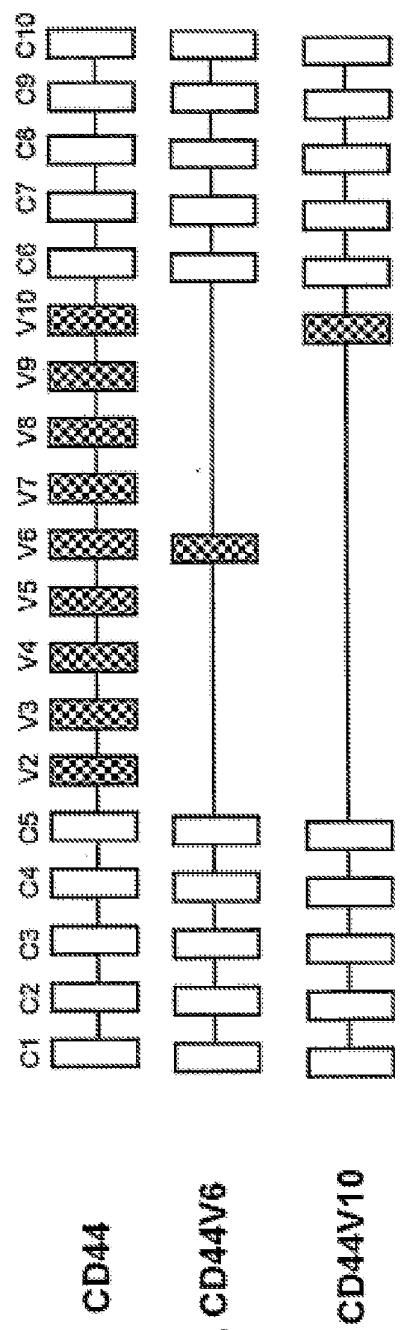


FIG. 13

PCTI L2012050104-seq1 -000001-EN
SEQUENCE LISTING

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<223> A pentapeptide derived from the human V6

<400> 1

Asn Arg Trp His Glu
1 5

<210> 2
<211> 39
<212> PRT
<213> artificial sequence

<220>
<223> Mouse V6 peptide

<400> 2

Pro Asn Ser Thr Ala Glu Ala Ala Ala Thr Glu Glu Glu Thr Trp Phe
1 5 10 15

Glu Asn Gly Trp Glu Gly Lys Asn Pro Pro Thr Pro Ser Glu Asp Ser
20 25 30

His Val Thr Glu Gly Thr Thr
35

<210> 3
<211> 20
<212> PRT
<213> artificial sequence

<220>
<223> Mouse V6A peptide

<400> 3

Pro Asn Ser Thr Ala Glu Ala Ala Ala Thr Glu Glu Glu Thr Trp Phe
1 5 10 15

Glu Asn Gly Trp

<210> 4
<211> 13
<212> PRT
<213> artificial sequence

<220>
<223> Mouse V6A1 peptide

<400> 4

Pro Asn Ser Thr Ala Glu Ala Ala Ala Thr Glu Glu Glu
1 5 10

<210> 5
<211> 6
<212> PRT
<213> artificial sequence

<220>
<223> Mouse V6A2 peptide

<400> 5

Trp Phe Glu Asn Gly Trp
1 5

<210> 6
<211> 12
<212> PRT
<213> artificial sequence

<220>
<223> Mouse V6B peptide

<400> 6

Thr Pro Ser Glu Asp Ser His Val Thr Glu Gly Thr
1 5 10

<210> 7
<211> 7
<212> PRT
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<220>
<223> Mouse V6B1 peptide

<400> 7

Thr Pro Ser Glu Asp Ser His
1 5

<210> 8
<211> 61
<212> PRT
<213> artificial sequence

<220>
<223> Human V10 peptide

<400> 8

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Thr Thr Leu Leu Glu Gly Tyr Thr Ser His Tyr Pro His Thr Lys Glu
1 5 10 15

Ser Arg Thr Phe Ile Pro Val Thr Ser Ala Lys Thr Gly Ser Phe Gly
20 25 30

Val Thr Ala Val Thr Val Gly Asp Ser Asn Ser Asn Val Asn Arg Ser
35 40 45

Leu Ser Gly Asp Gln Asp Thr Phe His Pro Ser Gly Gly
50 55 60

<210> 9

<211> 28

<212> PRT

<213> artificial sequence

<220>

<223> Human V10A peptide

<400> 9

Thr Thr Leu Leu Glu Gly Tyr Thr Ser His Tyr Pro His Thr Lys Glu
1 5 10 15

Ser Arg Thr Phe Ile Pro Val Thr Ser Ala Lys Thr
20 25

<210> 10

<211> 33

<212> PRT

<213> artificial sequence

<220>

<223> Human V10B peptide

<400> 10

Gly Ser Phe Gly Val Thr Ala Val Thr Val Gly Asp Ser Asn Ser Asn
1 5 10 15

Val Asn Arg Ser Leu Ser Gly Asp Gln Asp Thr Phe His Pro Ser Gly
20 25 30

Gly

<210> 11

<211> 16

<212> PRT

<213> artificial sequence

<220>

<223> Human V10A1 peptide

<400> 11

Thr Thr Leu Leu Glu Gly Tyr Thr Ser His Tyr Pro His Thr Lys Glu
Page 3

1

5

10

15

<210> 12
<211> 15
<212> PRT
<213> artificial sequence

<220>
<223> Human V10A2 peptide

<400> 12

Thr Lys Glu Ser Arg Thr Phe Ile Pro Val Thr Ser Ala Lys Thr
1 5 10 15

<210> 13
<211> 18
<212> PRT
<213> artificial sequence

<220>
<223> Human V10B1 peptide

<400> 13

Gly Ser Phe Gly Val Thr Ala Val Thr Val Gly Asp Ser Asn Ser Asn
1 5 10 15

Val Asn

<210> 14
<211> 21
<212> PRT
<213> artificial sequence

<220>
<223> Human V10B2 peptide

<400> 14

Asp Ser Asn Ser Asn Val Asn Arg Ser Leu Ser Gly Asp Gln Asp Thr
1 5 10 15

Phe His Pro Ser Gly
20

<210> 15
<211> 13
<212> PRT
<213> artificial sequence

<220>
<223> Human V10B3 peptide

<400> 15

Ser Leu Ser Gly Asp Gln Asp Thr Phe His Pro Ser Gly
1 5 10

<210> 16

<211> 7
<212> PRT
<213> Artificial sequence

<220>
<223> Human V6B1 peptide

<400> 16

Thr Pro Lys Glu Asp Ser His
1 5

<210> 17
<211> 4
<212> PRT
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<220>
<223> Human V6B1_C4 peptide

<400> 17

Gl u Asp Ser His
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<210> 18
<211> 17
<212> PRT
<213> Artificial sequence

<220>
<223> Human V10A/B_L peptide

<400> 18

Pro Val Thr Ser Ala Lys Thr Gly Ser Phe Gly Val Thr Ala Val Thr
1 5 10 15

Val

<210> 19
<211> 11
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<223> Human V10A/B_S peptide

<400> 19

Pro Val Thr Ser Ala Lys Thr Gly Ser Phe Gly
1 5 10

<210> 20
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<223> Human V10A2_C7 peptide

<400> 20

Pro Val Thr Ser Ala Lys Thr
1 5

<210> 21
<211> 10
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<220>
<223> Human V10A2_C10 peptide

<400> 21

Thr Phe Ile Pro Val Thr Ser Ala Lys Thr
1 5 10

<210> 22
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<220>
<223> Human V10A1_N12 peptide

<400> 22

Thr Thr Leu Leu Glu Gly Tyr Thr Ser His Tyr Pro
1 5 10

<210> 23
<211> 8
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<220>
<223> Human V10A1_N8 peptide

<400> 23

Thr Thr Leu Leu Glu Gly Tyr Thr
1 5

<210> 24
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<212> PRT
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<220>
<223> Human V10A1_N6 peptide

<400> 24

Thr Thr Leu Leu Glu Gly
1 5

<210> 25
<211> 6
<212> PRT
<213> Artificial sequence

<220>
<223> Human V10A1_N+2_6 peptide

<400> 25

Leu Leu Glu Glu Tyr Thr
1 5

<210> 26

<211> 8

<212> PRT

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<220>

<223> Human V10A1_N+4 peptide (also referred to as P26)

<400> 26

Gl u Gl y Tyr Thr Ser His Tyr Pro
1 5

<210> 27

<211> 4

<212> PRT

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<223> Human V10A1_EGYT peptide

<400> 27

Gl u Gl y Tyr Thr
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<210> 28

<211> 8

<212> PRT

<213> Artificial sequence

<220>

<223> Human V10B3_N7 peptide

<400> 28

Ser Leu Ser Gl y Asp Gl n Asp Thr
1 5

<210> 29

<211> 7

<212> PRT

<213> Artificial sequence

<220>

<223> Human V10B3_N6 peptide

<400> 29

Ser Leu Ser Gl y Asp Gl n Asp
1 5

<210> 30

<211> 5

<212> PRT

<213> Artificial sequence

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<220>
<223> Human V10B3_N5 peptide

<400> 30

Ser Leu Ser Gly Asp
1 5

<210> 31
<211> 6
<212> PRT
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<220>
<223> Human V10B2.5_6 peptide

<400> 31

Asn Val Asn Arg Ser Leu
1 5

<210> 32
<211> 9
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<220>
<223> Human V10B2.5_9 peptide

<400> 32

Asn Val Asn Arg Ser Leu Ser Gly Asp
1 5

<210> 33
<211> 10
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<220>
<223> Human V10B2_N10 peptide

<400> 33

Asp Ser Asn Ser Asn Val Asn Arg Ser Leu
1 5 10

<210> 34
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<223> Human V10B1_8 peptide (also referred to as P34)

<400> 34

Phe Gly Val Thr Ala Val Thr Val
1 5

<210> 35
<211> 7
<212> PRT

<213> Artificial sequence

<220>

<223> Human V10B1_7 peptide

<400> 35

Phe Gl y Val Thr Al a Val Thr
1 5

<210> 36

<211> 7

<212> PRT

<213> Artificial sequence

<220>

<223> Human 26-1 peptide

<400> 36

Gl u Gl y Tyr Thr Ser His Tyr
1 5

<210> 37

<211> 6

<212> PRT

<213> Artificial sequence

<220>

<223> Human 26-2 peptide

<400> 37

Gl u Gl y Tyr Thr Ser His
1 5

<210> 38

<211> 5

<212> PRT

<213> Artificial sequence

<220>

<223> Human 26-3 peptide

<400> 38

Gl u Gl y Tyr Thr Ser
1 5

<210> 39

<211> 7

<212> PRT

<213> Artificial sequence

<220>

<223> Human 26-4 peptide

<400> 39

Gl y Tyr Thr Ser His Tyr Pro
1 5

<210> 40

<211> 6
<212> PRT
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<220>
<223> Human 26-5 peptide

<400> 40

Tyr Thr Ser His Tyr Pro
1 5

<210> 41
<211> 5
<212> PRT
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<220>
<223> Human 26-6 peptide

<400> 41

Thr Ser His Tyr Pro
1 5

<210> 42
<211> 6
<212> PRT
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<220>
<223> Human 26-7 peptide

<400> 42

Gly Tyr Thr Ser His Tyr
1 5

<210> 43
<211> 8
<212> PRT
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<223> Human 26-8 peptide

<400> 43

Gln Gly Tyr Thr Ser His Tyr Pro
1 5

<210> 44
<211> 8
<212> PRT
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<220>
<223> Human 26-9 peptide

<400> 44

Glu Gly Tyr Thr Ser Ala Tyr Pro
1 5

<210> 45
<211> 8
<212> PRT
<213> Artificial sequence

<220>
<223> Human 26-RI peptide

<220>
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<223> D stereoisomer

<400> 45

Pro Tyr His Ser Thr Tyr Gly Glu
1 5

<210> 46
<211> 8
<212> PRT
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<220>
<223> Human 26-R peptide

<400> 46

Pro Tyr His Ser Thr Tyr Gly Glu
1 5

<210> 47
<211> 5
<212> PRT
<213> artificial sequence

<220>
<223> A minimal peptide sequence for mediating neuroprotection

<220>
<221> misc_feature
<222> (1)..(5)
<223> Can be D or L stereoisomer

<220>
<221> misc_feature
<222> (5)..(5)
<223> X can be E or O

<400> 47

Ser Thr Tyr Gly Xaa
1 5

<210> 48
<211> 8
<212> PRT
<213> artificial sequence

<220>
<223> A control peptide

<400> 48

Ala Val Ala Val Glu Ala Ala Gly
1 5

<210> 49

<211> 19

<212> PRT

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<220>

<223> An amino acid sequence missing from the CD44V10

<400> 49

Asp Ser Thr Asp Arg Ile Pro Ala Thr Ile Arg Asn Asp Val Thr Gly
1 5 10 15

Gly Arg Arg

<210> 50

<211> 19

<212> PRT

<213> artificial sequence

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<223> An amino acid sequence missing from the CD44V10

<400> 50

Asn Ser Asn Val Asn Arg Ser Leu Ser Gly Asp Gln Asp Thr Phe His
1 5 10 15

Pro Ser Gly

<210> 51

<211> 19

<212> PRT

<213> artificial sequence

<220>

<223> An amino acid sequence missing from the CD44 V6

<400> 51

Asp Ser Thr Asp Arg Ile Pro Ala Thr Ile Gln Ala Thr Pro Ser Ser
1 5 10 15

Thr Thr Glu

<210> 52

<211> 19

<212> PRT

<213> artificial sequence

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<223> An amino acid sequence missing from the CD44 V6

<400> 52

Asp	Ser	His	Ser	Thr	Thr	Gly	Thr	Ala	Gly	Asp	Gln	Asp	Thr	Phe	His
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Pro Ser Gly

<210> 53

<211> 742

<212> PRT

<213> Homo sapiens

<400> 53

Met	Asp	Lys	Phe	Trp	Trp	His	Ala	Ala	Trp	Gly	Leu	Cys	Leu	Val	Pro
1			5				10						15		

Leu	Ser	Leu	Ala	Gln	Ile	Asp	Leu	Asn	Ile	Thr	Cys	Arg	Phe	Ala	Gly
				20				25				30			

Val	Phe	His	Val	Glut	Lys	Asn	Gly	Arg	Tyr	Ser	Ile	Ser	Arg	Thr	Glut
				35			40				45				

Ala	Ala	Asp	Leu	Cys	Lys	Ala	Phe	Asn	Ser	Thr	Leu	Pro	Thr	Met	Ala
50				55						60					

Gln	Met	Glut	Lys	Ala	Leu	Ser	Ile	Gly	Phe	Glut	Thr	Cys	Arg	Tyr	Gly
65				70				75					80		

Phe	Ile	Glut	Gly	His	Val	Val	Ile	Pro	Arg	Ile	His	Pro	Asn	Ser	Ile
				85				90					95		

Cys	Ala	Ala	Asn	Asn	Thr	Gly	Val	Tyr	Ile	Leu	Thr	Ser	Asn	Thr	Ser
				100				105				110			

Gln	Tyr	Asp	Thr	Tyr	Cys	Phe	Asn	Ala	Ser	Ala	Pro	Pro	Glut	Glut	Asp
						115						120		125	

Cys	Thr	Ser	Val	Thr	Asp	Leu	Pro	Asn	Ala	Phe	Asp	Gly	Pro	Ile	Thr
130						135					140				

Ile	Thr	Ile	Val	Asn	Arg	Asp	Gly	Thr	Arg	Tyr	Val	Gln	Lys	Gly	Glut
145					150				155				160		

Tyr	Arg	Thr	Asn	Pro	Glut	Asp	Ile	Tyr	Pro	Ser	Asn	Pro	Thr	Asp	Asp
					165			170					175		

Asp	Val	Ser	Ser	Gly	Ser	Ser	Glu	Arg	Ser	Ser	Thr	Ser	Gly	Gly	
							180					185			

Tyr	Ile	Phe	Tyr	Thr	Phe	Ser	Thr	Val	His	Pro	Ile	Pro	Asp	Glut	Asp
								200				205			

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Ser Pro Trp Ile Thr Asp Ser Thr Asp Arg Ile Pro Ala Thr Thr Leu
210 215 220

Met Ser Thr Ser Ala Thr Ala Thr Glu Thr Ala Thr Lys Arg Glu Glu
225 230 235 240

Thr Trp Asp Trp Phe Ser Trp Leu Phe Leu Pro Ser Glu Ser Lys Asn
245 250 255

His Leu His Thr Thr Thr Glu Met Ala Glu Thr Ser Ser Asn Thr Ile
260 265 270

Ser Ala Gly Trp Glu Pro Asn Glu Glu Asn Glu Asp Glu Arg Asp Arg
275 280 285

His Leu Ser Phe Ser Gly Ser Gly Ile Asp Asp Asp Glu Asp Phe Ile
290 295 300

Ser Ser Thr Ile Ser Thr Thr Pro Arg Ala Phe Asp His Thr Lys Glu
305 310 315 320

Asn Glu Asp Trp Thr Glu Trp Asn Pro Ser His Ser Asn Pro Glu Val
325 330 335

Leu Leu Glu Thr Thr Arg Met Thr Asp Val Asp Arg Asn Glu Thr
340 345 350

Thr Ala Tyr Glu Gly Asn Trp Asn Pro Glu Ala His Pro Pro Leu Ile
355 360 365

His His Glu His His Glu Glu Glu Thr Pro His Ser Thr Ser Thr
370 375 380

Ile Glu Ala Thr Pro Ser Ser Thr Thr Glu Glu Thr Ala Thr Glu Lys
385 390 395 400

Glu Glu Trp Phe Gly Asn Arg Trp His Glu Glu Tyr Arg Glu Thr Pro
405 410 415

Lys Glu Asp Ser His Ser Thr Thr Glu Thr Ala Ala Ala Ser Ala His
420 425 430

Thr Ser His Pro Met Glu Glu Arg Thr Thr Pro Ser Pro Glu Asp Ser
435 440 445

Ser Trp Thr Asp Phe Phe Asn Pro Ile Ser His Pro Met Glu Arg Glu
450 455 460

His Glu Ala Glu Arg Arg Met Asp Met Asp Ser Ser His Ser Ile Thr
465 470 475 480

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Leu Glu Pro Thr Ala Asn Pro Asn Thr Gly Leu Val Glu Asp Leu Asp
485 490 495

Arg Thr Gly Pro Leu Ser Met Thr Thr Glu Glu Ser Asn Ser Glu Ser
500 505 510

Phe Ser Thr Ser His Glu Gly Leu Glu Glu Asp Lys Asp His Pro Thr
515 520 525

Thr Ser Thr Leu Thr Ser Ser Asn Arg Asn Asp Val Thr Gly Gly Arg
530 535 540

Arg Asp Pro Asn His Ser Glu Gly Ser Thr Thr Leu Leu Glu Glu Tyr
545 550 555 560

Thr Ser His Tyr Pro His Thr Lys Glu Ser Arg Thr Phe Ile Pro Val
565 570 575

Thr Ser Ala Lys Thr Gly Ser Phe Gly Val Thr Ala Val Thr Val Gly
580 585 590

Asp Ser Asn Ser Asn Val Asn Arg Ser Leu Ser Gly Asp Glu Asp Thr
595 600 605

Phe His Pro Ser Gly Gly Ser His Thr Thr His Gly Ser Glu Ser Asp
610 615 620

Gly His Ser His Gly Ser Glu Glu Gly Ala Asn Thr Thr Ser Gly
625 630 635 640

Pro Ile Arg Thr Pro Glu Ile Pro Glu Trp Leu Ile Ile Leu Ala Ser
645 650 655

Leu Leu Ala Leu Ala Leu Ile Leu Ala Val Cys Ile Ala Val Asn Ser
660 665 670

Arg Arg Arg Cys Gly Glu Lys Lys Lys Leu Val Ile Asn Ser Gly Asn
675 680 685

Gly Ala Val Glu Asp Arg Lys Pro Ser Gly Leu Asn Gly Glu Ala Ser
690 695 700

Lys Ser Glu Glu Met Val His Leu Val Asn Lys Glu Ser Ser Glu Thr
705 710 715 720

Pro Asp Glu Phe Met Thr Ala Asp Glu Thr Arg Asn Leu Glu Asn Val
725 730 735

Asp Met Lys Ile Gly Val
740

<210> 54
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<212> PRT
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<220>
<223> HIV TAT protein PTD derived cell penetrating peptide
<400> 54

Tyr Gl y Arg Lys Lys Arg Arg
1 5

<210> 55
<211> 11
<212> PRT
<213> artificial sequence

<220>
<223> HIV TAT protein PTD derived cell penetrating peptide
<400> 55

Tyr Gl y Arg Lys Lys Arg Arg Gl n Arg Arg Arg
1 5 10

<210> 56
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<212> PRT
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<220>
<223> HIV TAT protein PTD derived cell penetrating peptide
<400> 56

Arg Arg Gl n Arg Arg
1 5